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=> d his
     (FILE 'HOME' ENTERED AT 07:19:30 ON 01 FEB 2009)
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             1 S US20060204848/PN
               SAV L1 TEMP MAR947/A
               SEL L1 RN
    FILE 'REGISTRY' ENTERED AT 07:20:56 ON 01 FEB 2009
L2
            12 S E1-E12
               SAV TEMP L2 MAR947A/A
    FILE 'REGISTRY' ENTERED AT 07:33:49 ON 01 FEB 2009
L3
        460904 S (LI/ELS OR LITHIUM OR 7439-93-2/CRN) OR (NA/ELS OR SODI
L4
        1638539 S (MN/ELS OR MANGANESE OR 7439-96-5/CRN) OR (FE/ELS OR IR
    FILE 'LREGISTRY' ENTERED AT 07:38:47 ON 01 FEB 2009
L5
               OUE (SI/ELS OR SILICON OR 7440-21-3/CRN) OR (S/ELS OR SUL
    FILE 'REGISTRY' ENTERED AT 07:43:05 ON 01 FEB 2009
        1685641 S (P/ELS OR PHOSPHORUS OR 7723-14-0/CRN)
L6
L7
               QUE (O/ELS OR OXYGEN OR 17778-80-2/CRN)
T.8
          9432 S L3 AND L4 AND L5 AND L7
L9
          3503 S L3 AND L4 AND L6 AND L7
T-10
          2053 S L8 AND 4/0
L11
          1999 S L9 AND 4/0
               SAV TEMP L10 MAR947B/A
               SAV TEMP L11 MAR947C/A
L12
             1 S 33943-39-4
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L13
          2359 S L10
L14
          2864 S L11
L15
            98 S L12
L16
            55 S (LI# OR LITHIUM#) (W) (HYDROGEN#) (W) (PHOSPHATE#) OR L
L17
           131 S L15 OR L16
L18
             3 S L13 AND L17
L19
           15 S L14 AND L17
L20
           12 S L19 NOT L18
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FILE 'REGISTRY' ENTERED AT 08:11:07 ON 01 FEB 2009

1 S 67-42-5

L21

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             1 S 139-13-9
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L23
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          6823 S L22
L25
             1 S L13 AND L23
L26
             1 S L14 AND L23
L27
             1 S L13 AND L24
L28
             5 S L14 AND L24
T-29
             6 S L25-L28
L30
             5 S L29 NOT (L18 OR L20)
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                E DODECANE/CN
T.31
             1 S E3
               E TRIBUTYL PHOSPHATE/CN
L32
              1 S E3
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         12109 S L32
L33
L34
         12083 S L31
L35
             9 S (L10 OR L11) AND (L33 OR L34)
L36
             9 S L35 NOT (L18 OR L20 OR L30)
=> d his ful
     (FILE 'HOME' ENTERED AT 07:19:30 ON 01 FEB 2009)
     FILE 'HCAPLUS' ENTERED AT 07:19:53 ON 01 FEB 2009
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L1
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                SAV L1 TEMP MAR947/A
                SEL L1 RN
    FILE 'REGISTRY' ENTERED AT 07:20:56 ON 01 FEB 2009
L2
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                BI OR 139-13-9/BI OR 15365-14-7/BI OR 16448-54-7/BI OR
                21324-40-3/BI OR 24937-79-9/BI OR 33943-39-4/BI OR
                616-38-6/BI OR 67-42-5/BI OR 9003-07-0/BI OR 96-49-1/BI)
               D SCA
                SAV TEMP L2 MAR947A/A
     FILE 'STNGUIDE' ENTERED AT 07:23:32 ON 01 FEB 2009
     FILE 'REGISTRY' ENTERED AT 07:33:49 ON 01 FEB 2009
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L4 1638539 SEA SPE=ON ABB=ON PLU=ON (MN/ELS OR MANGANESE OR
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               (NI/ELS OR NICKEL OR 7440-02-0/CRN) OR (CO/ELS OR COBALT
               OR 7440-48-4/CRN)
   FILE 'LREGISTRY' ENTERED AT 07:38:47 ON 01 FEB 2009
T.5
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               (AL/ELS OR ALUMINUM OR 7429-90-5/CRN) OR (GE/ELS OR
               GERMANIUM OR 7440-56-4/CRN) OR (AS/ELS OR ARSENIC OR
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1.6
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               7723-14-0/CRN)
L7
               QUE SPE=ON ABB=ON PLU=ON (O/ELS OR OXYGEN OR 17778-80-
               2/CRN)
L8
          9432 SEA SPE=ON ABB=ON PLU=ON L3 AND L4 AND L5 AND L7
L9
          3503 SEA SPE=ON ABB=ON PLU=ON L3 AND L4 AND L6 AND L7
L10
         2053 SEA SPE=ON ABB=ON PLU=ON L8 AND 4/O
L11
          1999 SEA SPE=ON ABB=ON PLU=ON L9 AND 4/O
               SAV TEMP L10 MAR947B/A
              SAV TEMP L11 MAR947C/A
L12
             1 SEA SPE=ON ABB=ON PLU=ON 33943-39-4
    FILE 'HCAPLUS' ENTERED AT 07:51:38 ON 01 FEB 2009
         2359 SEA SPE=ON ABB=ON PLU=ON L10
L13
L14
          2864 SEA SPE=ON ABB=ON PLU=ON L11
L15
           98 SEA SPE=ON ABB=ON PLU=ON L12
L16
           55 SEA SPE=ON ABB=ON PLU=ON (LI# OR LITHIUM#) (W)
               (HYDROGEN#) (W) (PHOSPHATE#) OR LI2HPO4
L17
          131 SEA SPE=ON ABB=ON PLU=ON L15 OR L16
            3 SEA SPE=ON ABB=ON PLU=ON L13 AND L17
L18
L19
           15 SEA SPE=ON ABB=ON PLU=ON L14 AND L17
L20
            12 SEA SPE=ON ABB=ON PLU=ON L19 NOT L18
               D L20 HITSTR
    FILE 'STNGUIDE' ENTERED AT 08:03:07 ON 01 FEB 2009
    FILE 'REGISTRY' ENTERED AT 08:11:07 ON 01 FEB 2009
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L22
            1 SEA SPE=ON ABB=ON PLU=ON 139-13-9
    FILE 'HCAPLUS' ENTERED AT 08:11:38 ON 01 FEB 2009
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L23
          2855 SEA SPE=ON ABB=ON PLU=ON L21
L24
          6823 SEA SPE=ON ABB=ON PLU=ON L22
L25
             1 SEA SPE=ON ABB=ON PLU=ON L13 AND L23
L26
             1 SEA SPE=ON ABB=ON PLU=ON L14 AND L23
             1 SEA SPE=ON ABB=ON PLU=ON L13 AND L24
L27
L28
             5 SEA SPE=ON ABB=ON PLU=ON L14 AND L24
L29
             6 SEA SPE=ON ABB=ON PLU=ON (L25 OR L26 OR L27 OR L28)
             5 SEA SPE=ON ABB=ON PLU=ON L29 NOT (L18 OR L20)
L30
    FILE 'LREGISTRY' ENTERED AT 08:17:56 ON 01 FEB 2009
    FILE 'REGISTRY' ENTERED AT 08:20:48 ON 01 FEB 2009
               E DODECANE/CN
T.31
             1 SEA SPE=ON ABB=ON PLU=ON DODECANE/CN
               E TRIBUTYL PHOSPHATE/CN
1.32
             1 SEA SPE=ON ABB=ON PLU=ON "TRIBUTYL PHOSPHATE"/CN
    FILE 'HCAPLUS' ENTERED AT 08:21:18 ON 01 FEB 2009
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L33
L34
         12083 SEA SPE=ON ABB=ON PLU=ON L31
             9 SEA SPE=ON ABB=ON PLU=ON (L10 OR L11) AND (L33 OR
L35
               L34)
L36
             9 SEA SPE=ON ABB=ON PLU=ON L35 NOT (L18 OR L20 OR L30)
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FILE 'STNGUIDE' ENTERED AT 08:23:40 ON 01 FEB 2009

#### FILE HOME

### FILE HCAPLUS

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FILE COVERS 1907 - 1 Feb 2009 VOL 150 ISS 6
FILE LAST UPDATED: 29 Jan 2009 (20090129/ED)
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m HCAplus}$  now includes complete International Patent Classification (I reclassification data for the third quarter of 2008.

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DICTIONARY FILE UPDATES: 29 JAN 2009 HIGHEST RN 1097778-52-3

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## http://www.cas.org/support/stngen/stndoc/properties.html

#### FILE STUGUIDE

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LAST RELOADED: Jan 30, 2009 (20090130/UP).

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NEW CAS INFORMATION USE POLICIES, ENTER HELP USAGETERMS FOR DETAILS.

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YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L18 ANSWER 1 OF 3 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2008:816094 HCAPLUS Full-text

DN 149:204396

TI Preparation of metal-doped ferrous oxalate dihydrate as iron source

material for preparing metal-doped lithium iron(II) phosphate for use in lithium ion batteries

IN Cao, Wenyu; Zhang, Shuiyuan; Xiao, Feng

PA BYD Company Limited, Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 26pp.

CODEN: CNXXEV

DT Patent LA Chinese

LA Chinese FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	CN 101209820	A	20080702	CN 2006-10167328	200612

PRAI CN 2006-10167328

20061227

Metal-doped ferrous oxalate dihydrate is prepared by contacting a ferrous salt (ferrous sulfate, ferrous chloride and/or ferrous acetate) and a soluble nonferrous metal salt with an oxalate salt till the pH of the mixed solution is 3-6. The nonferrous metal salt can be a sulfate, nitrate and/or chloride of a IIA metal, IIIA metal, IVA metal, such as magnesium sulfate, aluminum sulfate, or zirconium sulfate. The oxalate can be sodium oxalate, potassium oxalate, ammonium oxalate, and/or lithium oxalate. The lithium iron phosphate is prepared by sintering a mixture of a lithium source, phosphorus source and the iron source material at 650-850° for 8-40 h in an inert gas or reducing gas atm; followed by cooling. The lithium source can be lithium hydroxide, lithium carbonate, or lithium acetate. The phosphorus source can be ammonium phosphate, ammonium hydrogen phosphate, or lithium phosphate. The mol. ratio of lithium to iron to phosphorus is (1-1.07):1:1. The obtained lithium iron(II) phosphate has a small particle size, uniform particles, good conductivity and electrochem, properties.

IT 554453-36-0P, Aluminum iron lithium phosphate

RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); PROC (Process); USES (Uses) (preparation of metal-doped ferrous oxalate dihydrate as iron

source

material for preparing metal-doped FeLiPO4 for use in lithium ion batteries)

RN 554453-36-0 HCAPLUS

CN Phosphoric acid, aluminum iron lithium salt (9CI) (CA INDEX NAME)

10/537 947

- ■x A1
- x Fe(x)
  - ●x Li

IT 33943-39-4, DiLithium hydrogen phosphate

RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent) (preparation of metal-doped ferrous oxalate dihydrate as iron

source

material for preparing metal-doped FeLiPO4 for use in lithium ion batteries)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

■ 2 T.i.

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49

IT 554453-36-0P, Aluminum iron lithium phosphate 554453-37-1P, Iron lithium zirconium phosphate 554453-38-2P, Iron lithium manganese phosphate 554453-42-8P, Iron lithium magnesium phosphate 912841-83-9P, Cobalt iron lithium phosphate 912841-84-0P, Iron lithium nickel phosphate

RL: IMF (Industrial manufacture); PEP (Physical, engineering or chemical process); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); PROC (Process); USES (Uses) (preparation of metal-doped ferrous oxalate dihydrate as iron

source

material for preparing metal-doped FeLiPO4 for use in lithium ion batteries)

ΙT 62-76-0, Sodium oxalate 546-89-4, Lithium acetate 553-91-3, Lithium oxalate 554-13-2, Lithium carbonate 583-52-8, Potassium oxalate 1113-38-8, Ammonium oxalate 1310-65-2, Lithium hydroxide 3094-87-9, Ferrous acetate 7446-70-0, Aluminum chloride, reactions 7646-79-9, Cobaltous chloride, reactions 7718-54-9, Nickelous chloride, reactions 7720-78-7, Ferrous sulfate 7722-76-1, Ammonium dihydrogen phosphate 7733-02-0, Zinc sulfate 7758-94-3, 7772-99-8, Stannous chloride, reactions Ferrous chloride 7773-01-5, Manganous chloride 7782-63-0, Ferrous sulfate heptahydrate 7783-28-0, Ammonium hydrogen phosphate 7784-31-8, Aluminum sulfate octadecahydrate 7785-87-7, Manganous sulfate 7786-30-3, Magnesium chloride, reactions 7790-69-4, Lithium nitrate 10034-99-8, Magnesium sulfate heptahydrate 10043-52-4, Calcium chloride, reactions 10099-59-9, Lanthanum nitrate 10101-97-0, Nickel sulfate hexahydrate 10124-43-3, Cobaltous 10361-37-2, Barium chloride, reactions 10361-65-6, TriAmmonium phosphate 10377-52-3, Lithium phosphate 10377-60-3, Magnesium nitrate 10476-85-4, Strontium chloride 13453-80-0, Lithium dihydrogen phosphate 13473-90-0, Aluminum nitrate 13746-89-9, Zirconium nitrate 33943-39-4, DiLithium hydrogen phosphate

RL: PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent)

(preparation of metal-doped ferrous oxalate dihydrate as iron

source

material for preparing metal-doped FeLiPO4 for use in lithium ion batteries)

L18 ANSWER 2 OF 3 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2005:409830 HCAPLUS Full-text

DN 142:466462

TI Product and method for the processing of precursors for lithium phosphate electrode active materials for batteries

IN Adamson, George; Barker, Jeremy; Dirilo, Allan; Faulkner, Titus; Saidi, Yazid M.; Swoyer, Jeffrey

PA Valence Technology, Inc., USA

SO PCT Int. Appl., 61 pp. CODEN: PIXXD2

DT Patent

LA English

WO 2004-US34229

FAN.		1 FENT	NO.			KIN		DATE			APPI	LICAT	ION :	NO.		D.	ATE
PI	WO	2005	- 0436	47				2005	0512		WO 2	2004-	US34	229		2	00410
	WO	2005	0436	47		A.3		2006	0511							-	_
		W:	CH, GB, KR, MX, SE,	CN, GD, KZ, MZ, SG,	CO, GE, LC, NA, SK,	CR, GH, LK, NI, SL,	CU, GM, LR, NO, SY,	CZ, HR, LS, NZ, TJ,	DE, HU, LT, OM,	DK, ID, LU, PG,	DM, IL, LV, PH,	BG, DZ, IN, MA, PL, TT,	EC, IS, MD, PT,	EE, JP, MG, RO,	EG, KE, MK, RU,	ES, KG, MN, SC,	FI, KP, MW, SD,
		RW:	BW, AM, DE, PT,	GH, AZ, DK, RO,	GM, BY, EE, SE,	KG, ES, SI,	LS, KZ, FI, SK,	MW, MD, FR,	RU, GB, BF,	TJ, GR,	TM,	SL, AT, IE, CG,	BE, IT,	BG, LU,	CH, MC,	CY, NL,	CZ, PL,
	US	2005									US 2	2004-	9616	73		2	00410
		7348 2542				B2 A1		2008 2005	0325 0512		CA 2	2004-	2542	790		2	00410
	DE	1120	0400	1997		Т5		2006	1026		DE 2	2004-	1120	0400	1997		00410
		1871										2004-				2	00410
		2008									US 2	2008-	4694	2		2	00803
PRAI	US US	2003 2004	-513: -961:	242P 673		P A		2003 2004	1021 1008								

AB The invention concerns methods for producing an electrode active material precursor, comprising: (a) producing a mixture comprising particles of lithium hydrogen phosphate, having a first average particle size, and a metal hydroxide, having a second average particle size; and (b) grinding the mixture in a jet mill for a period of time suitable to produce a generally homogeneous mixture of particles having a third average size smaller than the first average size. The precursor may be used as a starting material for making

20041015

W

electrode active materials for use in a battery, comprising lithium, a transition metal, and phosphate or a similar anion.

33943-39-4, DiLithium hydrogen phosphate

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)

(product and method for processing of precursors for lithium phosphate electrode active materials for batteries)

RN 33943-39-4 HCAPLUS

Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME) CN

610271-87-9 610271-90-4 610271-94-8 TT

610271-97-1 610272-06-5

RL: DEV (Device component use); USES (Uses)

(product and method for processing of precursors for lithium phosphate electrode active materials for batteries)

RN 610271-87-9 HCAPLUS

CN Aluminum cobalt iron lithium magnesium phosphate

(Al0.02Co0.85Fe0.05Li1.02Mg0.05(PO4)) (CA INDEX NAME)

Component	   	Ratio		Component Registry Number
			<b>-</b>	
04P	- 1	1	- 1	14265-44-2
Co	1	0.85	1	7440-48-4
Mg	1	0.05	- 1	7439-95-4
Li	1	1.02	- 1	7439-93-2
Fe	1	0.05	ĺ	7439-89-6
Al	1	0.02	1	7429-90-5

RN 610271-90-4 HCAPLUS

CN Aluminum cobalt iron lithium magnesium manganese phosphate (Al0.02Co0.7Fe0.08Li1.02Mg0.05Mn0.12(PO4)) (CA INDEX NAME)

Component Component | Registry Number

	====+====		+	
04P	1	1	1	14265-44-2
Co	1	0.7	1	7440-48-4
Mn	1	0.12	1	7439-96-5
Mg	1	0.05	1	7439-95-4
Li	1	1.02	1	7439-93-2
Fe	1	0.08	1	7439-89-6
A1	1	0.02	1	7429-90-5

RN 610271-94-8 HCAPLUS

CN Aluminum cobalt iron lithium magnesium phosphate (Al0.02Co0.8Fe0.1Li1.02Mg0.05(PO4)) (CA INDEX NAME)

Component	   	Ratio	Component   Registry Number
	-T		т
04P	1	1	14265-44-2
Co	1	0.8	7440-48-4
Mg	1	0.05	7439-95-4
Li	1	1.02	7439-93-2
Fe	1	0.1	7439-89-6
Al	1	0.02	7429-90-5

RN 610271-97-1 HCAPLUS

CN Aluminum cobalt iron lithium magnesium phosphate (Al0.02Co0.75Fe0.15Li1.02Mg0.05(PO4)) (CA INDEX NAME)

Component		Ratio		Component Registry Number
			т-	
04P	- 1	1	- 1	14265-44-2
Co	- 1	0.75	- 1	7440-48-4
Mg	1	0.05	- 1	7439-95-4
Li	1	1.02	- 1	7439-93-2
Fe	- 1	0.15	- 1	7439-89-6
Al	İ	0.02	ĺ	7429-90-5

RN 610272-06-5 HCAPLUS

CN Aluminum cobalt iron lithium titanium phosphate (Al0.02Co0.8Fe0.1Li1.02Ti0.02(PO4)) (CA INDEX NAME)

Component	1	Ratio	- 1	Component
	- 1		- 1	Registry Number
	+		===+==	
04P	1	1	- 1	14265-44-2
Co	- 1	0.8	- 1	7440-48-4
Ti	1	0.02	- 1	7440-32-6

Li	1	1.02	1	7439-93-2
Fe	1	0.1	1	7439-89-6
Al	1	0.02	1	7429-90-5

IC ICM H01M

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

ΙT 1305-62-0, Calcium hydroxide, processes 1309-37-1, Ferric oxide, 1309-42-8, Magnesium hydroxide 33943-39-4, processes

DiLithium hydrogen phosphate

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); PROC (Process)

(product and method for processing of precursors for lithium phosphate electrode active materials for batteries)

15365-14-7, Iron lithium phosphate felipo4 77641-62-4, Nasicon ΤТ 349632-85-5, Iron lithium magnesium phosphate Fe0.8LiMg0.2P04 610271-87-9 610271-90-4 610271-94-8

610271-97-1 610272-06-5 610310-97-9

610321-57-8 610321-60-3 610754-69-3 632286-77-2, 610321-55-6 Iron lithium magnesium phosphate Fe0.9LiMg0.1P04 771556-74-2

RL: DEV (Device component use): USES (Uses)

(product and method for processing of precursors for lithium phosphate electrode active materials for batteries)

RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L18 ANSWER 3 OF 3 HCAPLUS COPYRIGHT 2009 ACS on STN
- 2004:78425 HCAPLUS Full-text AN
- DN 140:131129
- Alkali metal hydrogen phosphates as precursors for electrode active TΤ materials
- TN Adamson, George W.; Barker, Jeremy; Faulkner, Titus; Saidi, M. Yazid: Swover, Jeffrev
- PA Valence Technology, Inc., USA
- SO U.S. Pat. Appl. Publ., 11 pp.
- CODEN: USXXCO
- DT Patent
- LA English

FAN.	CNT 1 PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	US 20040018135	A1	20040129	US 2002-205746	20020 <b>7</b> 26
	US 6794084 CA 2493224	B2 A1	20040921 20040205	CA 2003-2493224	

AB

IΤ

WO 2004011403 A2 20040205 WO 2003-US23167 200307 25 AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG CN 1672277 Α 20050921 CN 2003-817853 200307 25 CN 100334757 С 20070829 JP 2005533742 Т 20051110 JP 2004-524777 200307 25 PRAI US 2002-205746 20020726 Α WO 2003-US23167 W 20030725

An alkali metal hydrogen phosphate compound of the formula AxH3-xP04, wherein A is an alkali metal and 0<x<3, is prepared by mixing an alkali metal-containing basic compound, such as Li2CO3 or LiOH with phosphoric acid containing 5-25 weight% of water, agitating the mixture continuously, followed by drying and milling. An electrode active material is prepared having the general formula AaMb(PO4)cZd, wherein A is at least one alkali metal, M is at least one element capable of undergoing oxidation to a higher valence state, Z is a halogen or a hydroxy group, O<a\( 6 \), 1\le Ds3,

 $1 \le \le 3$ , and  $0 \le d \le 2$ , wherein M, Z, a, b, c and d are selected so as to maintain electroneutrality. To prepare the electrode active material an alkali metal hydrogen phosphate is mixed with an M-containing compound, followed by heating. M is at least one transition metal, such as Ti, V, Cr, Mn, Fe, Co, Ni, or Cu. The electrode active material can contain addnl. non-transition metals, such as Be, Mg, Ca, Sr, Ba Zn, Cd, Pb, Sn, Sc, Y, La, B, Al, Ga, or In. The manufacture electrodes the produced material is mixed with carbon. 649560-54-3P

RL: DEV (Device component use); NUU (Other use, unclassified); SPN (Synthetic preparation); PREP (Preparation); USES (Uses) (electrode material; alkali metal hydrogen phosphates as precursors for electrode active materials)

RN 649560-54-3 HCAPLUS

CN Aluminum cobalt iron lithium magnesium titanium phosphate (Al0.02Co0.8Fe0.1Li1.02Mg0.02Ti0.02(PO4)) (CA INDEX NAME)

Component		Ratio	   	Component Registry Number
			т	
04P	- 1	1		14265-44-2
Co	- 1	0.8		7440-48-4
Ti		0.02		7440-32-6
Mg		0.02		7439-95-4
Li	- 1	1.02	- 1	7439-93-2
Fe	- 1	0.1	- 1	7439-89-6
Al	- 1	0.02		7429-90-5

IC ICM C01B015-16

INCL 423313000

- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49
- ST lithium hydrogen phosphate electrode

material secondary battery

IT 649560-44-1P 649560-54-3P

RL: DEV (Device component use); NUU (Other use, unclassified); SPN (Synthetic preparation); PREP (Preparation); USES (Uses) (electrode material; alkali metal hydrogen phosphates as

(electrode material; alkali metal hydrogen phosphates as precursors for electrode active materials)

RE.CNT 14 THERE ARE 14 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

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YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

- L20 ANSWER 1 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN
- AN 2008:1487731 HCAPLUS Full-text

DN 150:80802

- TI Method for preparing lithium manganese phosphate as cathode material for lithium ion battery
- IN Yue, Min; Hou, Chunping; He, Xuegin; Zhang, Wanhong
- PA Shenzhen BTR New Energy Materials Inc., Peop. Rep. China
- SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 27pp. CODEN: CNXXEV
- DT Patent

LA Chinese

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	CN 101320809	A	20081210	CN 2008-10141632	
					200807

20080

PRAI CN 2008-10141632

20080717

AB The title cathode material is composed of lithium manganese phosphate particles coated with carbon material 1-3 weight% of lithium manganese phosphate. The cathode material has a sp. surface area of 5-40 m2/g and a tap d. of 1.0-1.6 g/mL. The title method comprises preparing nanoparticles, performing liquid-phase mixing reaction, preparing precursor, torrefying, and coating with the carbon material. The cathode material has high electronic conductivity, no agglomeration, high charge/discharge capacity, high cycle stability, high safety, easy preparation, low cost, and little influence to environment.

IT 13826-59-0P, Lithium manganese phosphate

RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(method for preparing lithium manganese phosphate as cathode material for lithium ion battery)

RN 13826-59-0 HCAPLUS

CN Phosphoric acid, lithium manganese(2+) salt (1:1:1) (9CI) (CA INDEX NAME)

■ Li

Mn(II)

IT 33943-39-4, Dilithium hydrogen phosphate
RL: RCT (Reactant); RACT (Reactant or reagent)
(method for preparing lithium manganese phosphate as cathode

material for lithium ion battery) 33943-39-4 HCAPLUS Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

RN

CN

●2 Li

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49

IT 13826-59-0P, Lithium manganese phosphate
RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or
engineered material use); PREP (Preparation); USES (Uses)
(method for preparing lithium manganese phosphate as cathode
material for lithium ion battery)

50-81-7, Ascorbic acid, reactions 50-99-7, Glucose, reactions ΙT 56-81-5, Glycerol, reactions 57-13-6, Urea, reactions 57-48-7, Fructose, reactions 57-50-1, Sucrose, reactions 77-92-9, Citric acid, reactions 79-10-7, Acrylic acid, reactions 87-69-4, Tartaric acid, reactions 546-89-4, Lithium acetate 553-54-8, Lithium benzoate 553-91-3, Lithium oxalate 554-13-2. Lithium carbonate 556-63-8, Lithium formate 919-16-4, Lithium citrate 1310-65-2, Lithium hydroxide 1313-13-9, Manganese dioxide, reactions 1314-56-3, Phosphorus pentoxide, reactions 1317-34-6, Manganese sesquioxide 1907-33-1 2180-18-9, Manganese acetate 7447-41-8, Lithium chloride, reactions 7550-35-8, Lithium bromide 7558-79-4, Disodium hydrogen phosphate 7558-80-7, Sodium dihydrogen phosphate 7664-38-2, Phosphoric acid, reactions 7722-76-1, Ammonium dihydrogen phosphate 7758-11-4, Dipotassium hydrogen phosphate 7778-77-0, Potassium dihydrogen phosphate 7783-28-0. Diammonium hydrogen phosphate 7789-24-4. Lithium fluoride, reactions 7790-69-4, Lithium nitrate 9002-84-0, Polytetrafluoroethylene 9002-89-5, Polyvinyl alcohol 9003-08-1, Melamine resin 9003-53-6, Polystyrene 9004-34-6, Cellulose, reactions 9011-05-6, Urea-formaldehyde resin 9011-14-7. Polymethyl methacrylate 10124-31-9, Ammonium phosphate 10377-48-7, Lithium sulfate 10377-51-2, Lithium iodide 10377-52-3, Lithium phosphate 12626-88-9, Manganese hydroxide 13453-80-0, Lithium dihydrogen phosphate 14024-11-4, Lithium

aluminum tetrachloride 14283-07-9, Lithium tetrafluoroborate 14307-35-8, Lithium chromate 24937-79-9, Poly(vinylidene fluoride) 25014-41-9, Polyacrylonitrile 25322-68-3, Polyethylene glycol

33943-39-4, Dilithium hydrogen phosphate

RL: RCT (Reactant); RACT (Reactant or reagent)

(method for preparing lithium manganese phosphate as cathode material for lithium ion battery)  $% \left( \frac{1}{2}\right) =\frac{1}{2}\left( \frac{1}{2}\right) +\frac{1}{2}\left( \frac{1}{2}\right) +\frac$ 

L20 ANSWER 2 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2008:875000 HCAPLUS Full-text

DN 149:248763

TI Method for preparing electrode material with ferrophosphorus

IN Wang, Guixin; Yan, Kangping

PA Sichuan University, Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 9pp.

DT Patent

LA Chinese FAN.CNT 1

PΙ

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 101219783	A	20080716	CN 2008-10045243	200801

23

PRAI CN 2008-10045243

20080123

AB The title method can prepare electrode material such as LiFePO4, LiFePO4/FeP2, LiFePO4/C, LiJFe2(PO4)3, FeP, FeP2, Fe2P, Fe3P, Fe-Co-P, Fe-Ni-P, Fe-Ni-Co-P, etc. from ferrophosphorus with or without addition of other elements by mech. activation method, reaction pulverization method, rheol. phase reaction method, spray drying method, spray pyrolysis method, solid phase method, microwave method, H2O/alc. thermal synthesis method, sol-gel method, ion exchange method, etc. The method has the advantages of wide raw material resources, low cost, simple operation, short flow process, etc., and realizes comprehensive use of resources.

II 33943-39-4, Dilithium hydrogen phosphate

RL: RCT (Reactant); RACT (Reactant or reagent)

(method for preparing electrode material with ferrophosphorus)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

■2 Ld

● Fe(II)

● Li

RN 36058-25-0 HCAPLUS
CN Phosphoric acid, iron(3+) lithium salt (3:2:3) (9CI) (CA INDEX NAME)

10/537,947

●2/3 Fe(III)

● Tri

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CC 49-5 (Industrial Inorganic Chemicals)
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Section cross-reference(s): 52

IT 546-89-4, Lithium acetate 554-13-2, Lithium carbonate 1310-65-2, Lithium hydroxide 10377-52-3, Lithium phosphate 13453-80-0, Lithium dihydrogen phosphate 33943-39-4, Dilithium hydrogen phosphate

RL: RCT (Reactant); RACT (Reactant or reagent)

(method for preparing electrode material with ferrophosphorus)

IT 12674-76-9P 15365-14-7P, Iron lithium phosphate (FeLiPO4)
36058-25-0P, Iron lithium phosphate (Fe2Li3(PO4)3)

50954-84-2P 71849-39-3P

RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (method for preparing electrode material with ferrophosphorus)

L20 ANSWER 3 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2008:735672 HCAPLUS Full-text

DN 149:152744

 ${\tt TI}$  Method for preparing ferrous oxalate used in preparation of lithium ferrous phosphate

IN Cao, Wenyu; Zhang, Shuiyuan; Xiao, Feng

PA Byd Co., Ltd., Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 18pp.

CODEN: CNXXEV
DT Patent

LA Chinese

FAN.CNT 1

PI CN 101200422 A 20080618 CN 2006-10167409

200612

15

PRAI CN 2006-10167409 20061215

AB The invention discloses a method for preparing ferrous oxalate through performing contact between ferrous salt solution flow and oxalate solution flow. The pH value of the obtained mixture is controlled at 2-6 by adjusting the flow rates of the ferrous salt solution flow and oxalate solution flow. By the method, lithium ferrous phosphate particles with high uniformity, small sizes, high carbon distribution uniformity, and good electrochem. properties can be obtained.

IT 15365-14-7P, Ferrous lithium phosphate

RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(method for preparing ferrous oxalate used in preparation of

lithium

ferrous phosphate)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

■ Li

IT 33943-39-4, Dilithium hydrogenphosphate

RL: RGT (Reagent); RACT (Reactant or reagent)

(method for preparing ferrous oxalate used in preparation of

lithium ferrous phosphate)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

10/537 947

■2 Td

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CC 23-16 (Aliphatic Compounds)
    Section cross-reference(s): 45, 72
ΙT
    15365-14-7P, Ferrous lithium phosphate
    RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP
     (Preparation)
        (method for preparing ferrous oxalate used in preparation of
lithium
       ferrous phosphate)
    546-89-4, Lithium acetate 554-13-2, Lithium carbonate 1310-65-2,
ΙT
    Lithium hydroxide 3094-87-9, Ferrous acetate 7720-78-7, Ferrous
    sulfate 7722-76-1, Ammonium dihydrogenphosphate 7758-94-3,
    Ferrous chloride 7783-28-0, Diammonium phosphate 7790-69-4,
    Lithium nitrate 10361-65-6, Triammonium phosphate 10377-52-3,
    Lithium phosphate 13453-80-0, Lithium dihydrogenphosphate
     33943-39-4, Dilithium hydrogenphosphate
     RL: RGT (Reagent); RACT (Reactant or reagent)
       (method for preparing ferrous oxalate used in preparation of
lithium
       ferrous phosphate)
L20 ANSWER 4 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN
AN
    2008:450742 HCAPLUS Full-text
DN
    148:520660
   LiFePO4/C nano-composite cathode material and its manufacture
TI
IN Xu, Yunlong; Ma, Hongyan; Tao, Lili
PA Shanghai Weina Company, Peop. Rep. China
SO Faming Zhuanli Shenging Gongkai Shuomingshu, 11pp.
    CODEN: CNXXEV
DT Patent
LA Chinese
FAN.CNT 1
    PATENT NO.
                 KIND DATE APPLICATION NO. DATE
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PI CN 101159328 A 20080409 CN 2007-10043889

200707

AB

20070717

17

PRAI CN 2007-10043889

The title cathode material is obtained by (1) weighing a Li source, an iron source, and a phosphorus source at a molar ratio of (3.0-3.3):(1.0-1.1):(1.0-1.1), and adding in a reaction container with an appropriate quantity of a carbon doped material and organic surfactant, (2) controlling the concentration and temperature of reaction solution to obtain a precursor gel, separating, washing, filtering and drying to obtain a precursor powder, and (3) tableting, putting in a crucible having a microwave absorbent, placing the crucible in a microwave oven, and heating for 3-30 min under 100-600 W to obtain the final product. The method has short preparation period, low energy consumption, and easy control of process, and is suitable for industrial production The cathode material has high purity, small particle size (< 100 nm), and good electrochem.

IT 15365-14-7P, Iron lithium phosphate (FeLiPO4)
RL: IMF (Industrial manufacture); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(manufacture of LiFePO4/C composite cathode materials for secondary

lithium batteries)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

IT 33943-39-4, Dilithium hydrogen phosphate

RL: RCT (Reactant); RACT (Reactant or reagent)

(manufacture of LiFePO4/C composite cathode materials for secondary

lithium batteries)

RN 33943-39-4 HCAPLUS
CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

но— Р— он

●2 Li

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT 15365-14-7P, Iron lithium phosphate (FeLiPO4)
RL: IMF (Industrial manufacture); TEM (Technical or engineered
material use); PREP (Preparation); USES (Uses)

 $\begin{tabular}{ll} (manufacture of LiFePO4/C composite cathode materials for secondary \end{tabular}$ 

lithium batteries)

IT 546-89-4, Lithium acetate 1310-65-2, Lithium hydroxide 7447-41-8, Lithium chloride, reactions 7664-38-2, Phosphoric acid, reactions 7720-78-7, Ferrous sulfate 7722-76-1, Ammonium dihydrogen phosphate 7783-28-0, Diammonium hydrogen phosphate 7790-69-4, Lithium nitrate 10045-89-3, Ammonium ferrous sulfate 10421-48-4, Ferric nitrate 13453-80-0, Lithium dihydrogen phosphate 33943-39-4, Dilithium hydrogen phosphate RL: RCT (Reactant); RACT (Reactant or reagent)

(manufacture of LiFePO4/C composite cathode materials for secondary

lithium batteries)

L20 ANSWER 5 OF 12 HCAPLUS COPYRIGHT 2009 ACS on SIN

AN 2008:428956 HCAPLUS Full-text

DN 148:474802

TI Preparation method of lithium iron phosphate used as cathode active material for lithium ion secondary battery

IN Liu. Fei

PA Byd Company Limited, Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 20pp. CODEN: CNXXEV

DT Patent

LA Chinese

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

CN 101152960 A 20080402 CN 2006-10152271 PI

> 200609 27

PRAI CN 2006-10152271

20060927

AB The title method comprises mixing elec. conductive particles, ferric ion- or ferrous ion-containing solution, and phosphate-containing solution at an Fe/P mol. ratio of (1-1.3):1, precipitating, separating solid, washing to obtain ferric or ferrous phosphate precipitation containing elec. conductive particles, mixing with Li source, and calcining at 500-900° for 8-48 h in inert or reducing atmospheric The cathode active material has good crystal structure and high specific capacitance.

15365-14-7P, Iron lithium phosphate (FeLiPO4) ΙT

RL: IMF (Industrial manufacture); PRP (Properties); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (preparation of lithium iron phosphate as cathode active material

for lithium ion secondary battery)

15365-14-7 HCAPLUS RN

> Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

CN

Fe(II)

Li

for

33943-39-4, Dilithium hydrogen phosphate TT

RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of lithium iron phosphate as cathode active material

lithium ion secondary battery)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

●2 Li

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CC 49-5 (Industrial Inorganic Chemicals)
Section cross-reference(s): 52
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for

lithium ion secondary battery)

T56-03-0, Ferrous oxalate 546-89-4, Lithium acetate 553-91-3, Lithium oxalate 554-13-2, Lithium carbonate 1310-65-2, Lithium hydroxide 2944-66-3, Ferric oxalate 7447-41-8, Lithium chloride, reactions 7664-38-2, Phosphoric acid, reactions 7705-08-0, Ferric chloride, reactions 7720-78-7, Ferrous sulfate 7722-76-1, Ammonium dihydrogen phosphate 7758-94-3, Ferrous chloride 7783-28-0, Diammonium hydrogen phosphate 7790-69-4, Lithium nitrate 10028-22-5, Ferric sulfate 10361-65-6, Ammonium phosphate 10377-52-3, Lithium phosphate 10421-48-4, Ferric nitrate 13453-80-0, Lithium dihydrogen phosphate 14013-86-6, Ferrous nitrate 33943-39-4, Dillthium hydrogen phosphate RL: RCT (Reactant); RACT (Reactant or reagent)

(preparation of lithium iron phosphate as cathode active material

for

lithium ion secondary battery)

- L20 ANSWER 6 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN
- AN 2008:39052 HCAPLUS Full-text
- DN 148:148439
- TI Preparation and application of LiFeP04/Li3V2(P04)3 composite cathode materials for lithium ion batteries
- IN Wu, She-Huang; Yang, Mu-Rong; Ke, Wei-Hsin; Huang, Yuan-Lung; Yu, Nien-Chieh
- PA Tatung Company, Taiwan
- SO U.S. Pat. Appl. Publ., 11pp. CODEN: USXXCO

DT Patent LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PΙ	US 20080008938	A1	20080110	US 2007-783299	
					200704

09

PRAI TW 2006-95124642 A 20060706

A method of preparing LiFePO4/Li3V2(PO4)3 composite cathode materials AB and their applications as cathode materials for lithium ion batteries are disclosed. The preparation method includes the following steps: (A) providing a mixture of iron powder, lithium salt, vanadium salt, and a phosphate salt whereafter these compds. are dissolved into a mixed acid solution; (B) drving the solution in order to obtain precursor powders; and (C) heating the precursor powders at a temperature ranging between 400 and 1000° to form LiFe1y'Vy'P04/Li3V2-y"Fey"(P04)3 composite powders. Alternatively, prepare the composite cathode by preparing olivine LiFe1-y'Vy'PO4 and monoclinic Li3V2-y'Fey"(PO4)3 powders as in previous procedures followed by mixing adequately. The low cost of iron powder thus facilitates to prepared composite cathode materials exhibiting higher elec. conductivity and superior cycling performance at high rates than those of olivine LiFel-y'Vy'PO4 and monoclinic Li3V2y"Fey" (PO4)3. The invention will help the development of the lithium ion batteries and related industries.

IT 33943-39-4, DiLithium hydrogen phosphate

RL: RCT (Reactant); RACT (Reactant or reagent) (preparation and application of LiFePO4/Li3V2(PO4)3 composite cathode

materials for lithium ion batteries)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

●2 Li

IT 15365-14-7P, Iron lithium phosphate felipo4
RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)
(preparation and application of LiFePO4/Li3V2(PO4)3 composi

(preparation and application of LiFePO4/Li3V2(PO4)3 composite cathode

materials for lithium ion batteries)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

• Fe(II)

● Li

INCL -429; -205; 252182100

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49

64-19-7, Acetic acid, reactions 77-92-9, Citric acid, reactions IΤ 79-09-4, Propionic acid, reactions 87-69-4, Tartaric acid, reactions 107-92-6, Butyric acid, reactions 144-62-7, Oxalic acid, reactions 546-89-4, Lithium acetate 1310-65-2, Lithium hydroxide (Li(OH)) 1314-34-7, Vanadium oxide (V2O3) 1314-56-3, Phosphorus oxide (P205), reactions 7439-89-6, Iron, reactions 7447-41-8, Lithium chloride, reactions 7550-35-8, Lithium bromide 7601-90-3, Perchloric acid, reactions 7647-01-0, Hydrochloric acid, reactions 7664-38-2, Phosphoric acid, reactions 7664-39-3, Hydrofluoric acid, reactions 7664-93-9, Sulfuric acid, reactions 7697-37-2, Nitric acid, reactions 7722-76-1, Ammonium dihydrogen phosphate 7789-24-4, Lithium fluoride, reactions 7790-69-4, Lithium nitrate 7790-92-3, Hypochlorous acid 7803-55-6, Ammonium vanadate 10361-65-6, Triammonium phosphate 10377-52-3, Lithium phosphate 12036-21-4, Vanadium oxide (VO2) 12057-24-8, Lithium oxide, reactions 13453-80-0, Lithium dihydrogen phosphate 33943-39-4, DiLithium hydrogen phosphate 64580-71-8 1001050-87-8

RL: RCT (Reactant); RACT (Reactant or reagent) (preparation and application of LiFePO4/Li3V2(PO4)3 composite cathode

materials for lithium ion batteries)

15365-14-7P. Iron lithium phosphate felipo4 84159-18-2P. IT

Lithium vanadium phosphateLi3V2(PO4)3

RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (preparation and application of LiFePO4/Li3V2(PO4)3 composite

cathode

materials for lithium ion batteries)

L20 ANSWER 7 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN

2007:850926 HCAPLUS Full-text AN

147:280811 DN

Method for preparing LiFePO4 particles with controllable morphology TT

IN Ni, Jiangfeng; Zhou, Henghui; Chen, Jitao; Zhang, Xinxiang

Pulead Technology Industry Co., Ltd., Peop. Rep. China PA

SO Faming Zhuanli Shenging Gongkai Shuomingshu, 12pp.

CODEN: CNXXEV

Patent. DT

FAN.	CNI 1				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PΤ	CN 101007630	A	20070801	CN 2007-10000683	
					200701 16

PRAI CN 2007-10000683 20070116

- The title method comprises the steps of: (1) mixing one or more kinds AB of compds. or solns. containing lithium ions, iron ions, and phosphate ions, adding solvent, adding crystal growth inhibitor (0.5-50 weight% of the theoretic product), and transferring to a hermetic reaction kettle. (2) performing solvent-thermal reaction to obtain the primary product, and (3) cooling, washing, filtering, and drying. The product can be calcined at high temperature for higher crystallinity. The LiFePO4 is useful as cathodic substance of lithium ion batteries for elec. tools, elec. bicycles, and elec. automobiles. The LiFePO4 particles have the advantages of various kinds of morphol., uniform size distribution, high controllability of morphol. and size, and small particles size. The method can be used for synthesizing submicroscale and nanoscale products, and has the advantages of short reaction time and low energy consumption.
- 15365-14-7P, Iron lithium phosphate, (LiFePO4) ΙT RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(method for preparing LiFePO4 particles with controllable morphol.)  $% \label{eq:controllable}% \end{substitute} % \begin{substitute}[t]{0.9\textwidth} \includegraphics[width=\linewidth]{images/controllable} \includegraphics[width=\li$ 

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

IT 33943-39-4, Dilithium hydrogen phosphate

RL: RCT (Reactant); RACT (Reactant or reagent)

(method for preparing LiFePO4 particles with controllable morphol.)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

■ 2 T 4

- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49
- IT 15365-14-7P, Iron lithium phosphate, (LiFePO4)
  RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)
  (method for preparing LiFePO4 particles with controllable

```
morphol.)

IT 546-89-4, Lithium acetate 1310-65-2, Lithium hydroxide
1332-37-2, Iron oxide, reactions 3094-87-9, Ferrous acetate
7664-38-2, Phosphoric acid, reactions 7720-78-7, Ferrous sulfate
7722-76-1, Ammonium dihydrogen phosphate 7758-94-3, Ferrous
chloride 7783-28-0, Diammonium hydrogen phosphate 10045-86-0,
Ferric phosphate 10124-31-9, Ammonium phosphate 10290-71-8, Iron
carbonate 10377-52-3, Lithium phosphate 11113-65-8, Iron
fluoride 11126-12-8, Iron sulfide 13453-80-0, Lithium dihydrogen
phosphate 14013-86-6, Iron nitrate 14940-41-1, Ferrous phosphate
33943-39-4, Dilithium hydrogen phosphate 52767-99-4,
Ammonium iron phosphate
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RL: RCT (Reactant); RACT (Reactant or reagent)

(method for preparing LiFePO4 particles with controllable morphol.)  $\,$ 

L20 ANSWER 8 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2006:1015282 HCAPLUS Full-text

DN 145:474775

TI Method for manufacturing lithium ferrous phosphate as cathode material of lithium-ion batteries

IN Gu, Yijie; Huang, Xiaowen; Cui, Hongzhi

PA Shandong University of Science and Technology, Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 5 pp. CODEN: CNXXEV

DT Patent LA Chinese

FAN.CNT 1

PΙ

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 1837033	A	20060927	CN 2006-10043350	200603

24

CN 100413781 C 20080827 PRAI CN 2006-10043350 20060324

AB The title method comprises: (1) mixing lithium salts, ferrous salts and ammonium dihydrogen phosphate at a mol. ratio (lithium ion to ferrous ion to phosphate radical) of (0.8-1.2):(0.8-1.2):(0.8-1.2) to obtain mixture A, (2) adding the mixture A in solution B (aqueous solution containing dissolvable salts and organic substances) at a weight ratio of 1:(0.1-10), stirring, placing into a high-temperature furnace, heating without air or oxidative gas atmospheric at a rate of 1-30°/min, keeping the temperature of 50-200° for 0-100 h (the higher the temperature is, the shorter the time is), carrying out high-temperature treatment by elec. heating, and cooling naturally to obtain lithium ferrous phosphate (LixFeyMzPO4) powder, and (3)

grinding the powder to a particle size of 1-50  $\mu m$  to obtain the final product. In step 1, lithium salt is one of lithium carbonate, lithium hydroxide, dilithium hydroxepen phosphate, lithium sulfate, lithium acetate, lithium nitrate and lithium oxalate, and ferrous salt is ferrous acetate or ferrous oxalate. In solution B, the dissolvable salt (M) is at least one of nitrate, acetate, sulfate, and chloride of aluminum, titanium, magnesium, zirconium, vanadium, manganese, nickel, cobalt, niobium, rhodium, barium, and chromium with a doping amount of M/lithium mol. ratio of  $\leq 0.3$ , and the dissolvable organic substance is at least one of sucrose, glucose, and macromol. compound pyrolyzed into carbon substances with good elec. conductivity with a doping amount of carbon/final product weight ratio  $\leq 10$ . The title cathode material has the advantages of uniform distribution, and improved charge capacity.

15365-14-7P, Ferrous lithium phosphate

RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP (Preparation)

(magnesium or zirconium doped; process for manufacturing ferrous lithium phosphate as cathode active material for lithium ion batteries)

RN 15365-14-7 HCAPLUS

Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

ΙT

CN

● Fe(II)

● Li

IT 33943-39-4, Dilithium hydrogen phosphate

RL: RCT (Reactant); RACT (Reactant or reagent)

(process for manufacturing ferrous lithium phosphate as cathode active

material for lithium ion batteries)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

10/537 947

●2 Li

- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49
- IT 15365-14-7F, Ferrous lithium phosphate
  RL: IMF (Industrial manufacture); SPN (Synthetic preparation); PREP
   (Preparation)

(magnesium or zirconium doped; process for manufacturing ferrous lithium phosphate as cathode active material for lithium ion

batteries)

IT 50-99-7, Glucose, reactions 57-50-1, Sucrose, reactions 516-03-0, Ferrous oxalate 546-89-4, Lithium acetate 553-91-3, Lithium oxalate 554-13-2, Lithium carbonate 1310-65-2, Lithium hydroxide 3094-87-9, Ferrous acetate 7722-76-1, Ammonium dihydrogen phosphate 7790-69-4, Lithium nitrate 10377-48-7, Lithium sulfate 33943-39-4, Dilithium hydrogen phosphate RL: RCT (Reactant); RACT (Reactant or reagent)

 $\hbox{(process for manufacturing ferrous lithium phosphate as cathode} \\$ 

material for lithium ion batteries)

L20 ANSWER 9 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2006:878134 HCAPLUS Full-text

DN 146:29497

TI Method for preparing spherical or quasi-spherical metal lithium phosphate

IN Ni, Jiangfeng; Zhou, Henghui; Chen, Jitao; Zhang, Xinxiang

PA Pulead Technology Industry Co., Ltd, Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 10pp.

CODEN: CNXXEV

DT Patent LA Chinese

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE

PI CN 1821063 A 20060823 CN 2006-10011378

200602 28

CN 100390052 C 20080528 PRAI CN 2006-10011378 20060228

AB The title method comprises: (1) pulverizing one or more compds. containing lithium ion, transition metal ions, and phosphate, (2) pyrolyzing under inert gas atmospheric, (3) adding molten alkali metal salts, wherein the mol. ratio of molten salts/transition metal ions is 0.1-10, and sintering, and (4) cooling, washing, filtering, drying, and pulverizing to obtain the final product with a particle size of 1-5 µm. The particle size of the final product can be controlled by reaction conditions. The method has the advantages of short sintering time requirement and low energy consumption. The obtained product has the advantages of 1 low sp. surface area, good processing property, high tap d., high volumetric specific energy d., and good safety. The product can be widely used in batteries of elect tools, elect bicycles, and elect cars.

IT 33943-39-4, Dilithium hydrogen phosphate

RL: PEP (Physical, engineering or chemical process); PROC (Process) (method for preparing spherical or quasi-spherical metal lithium phosphate)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

■2 T.i

IT 13826-59-0P, Lithium manganese phosphate 15365-14-7P, Ferrous lithium phosphate 153456-60-1P 554453-38-2P, Iron lithium manganese phosphate 915945-24-3P, Copper iron lithium zinc phosphate 915945-25-4P, Cobalt iron lithium manganese phosphate RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses) (method for preparing spherical or quasi-soherical metal lithium

phosphate)
RN 13826-59-0 HCAPLUS

CN Phosphoric acid, lithium manganese(2+) salt (1:1:1) (9CI) (CA INDEX NAME)

- Li
- Mn(II)
- RN 15365-14-7 HCAPLUS CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

- Fe(II)
  - Li

RN 153456-60-1 HCAPLUS CN Phosphoric acid, cobalt lithium nickel salt (9CI) (CA INDEX NAME) 10/537,947

- •x Co(x)
  - ●x Li
- ●x Ni(x)

RN 554453-38-2 HCAPLUS CN Phosphoric acid, iron lithium manganese(2+) salt (9CI) (CA INDEX NAME)

- ●x Fe(x)
  - ●x Li
- Mn(II)

RN 915945-24-3 HCAPLUS CN Copper iron lithium zinc phosphate (CA INDEX NAME)

Component | Ratio | Component | Registry Number

04P	1	х	1	14265-44-2
Zn	1	х	1	7440-66-6
Cu	1	х	1	7440-50-8
Li	1	х	1	7439-93-2
Fe	1	x	I	7439-89-6

RN 915945-25-4 HCAPLUS

CN Cobalt iron lithium manganese phosphate (CA INDEX NAME)

Component	 	Ratio	 	Component Registry Number
	+		+-	
04P	- 1	x	- 1	14265-44-2
Co	- 1	x	- 1	7440-48-4
Mn	- 1	x	1	7439-96-5
Li	- 1	x	- 1	7439-93-2
Fe	- 1	х	- 1	7439-89-6

CC 49-5 (Industrial Inorganic Chemicals)

- ΙT 142-71-2, Copper acetate 516-03-0, Ferrous oxalate 546-89-4, Lithium acetate 554-13-2, Lithium carbonate 1309-37-1, Ferric oxide, processes 1310-65-2, Lithium hydroxide 1313-99-1. Nickel oxide, processes 1317-37-9, Ferrous sulfide 2180-18-9, Manganese acetate 3486-35-9, Zinc carbonate 7542-09-8, Cobalt carbonate 7664-38-2, Phosphoric acid, processes 7722-76-1, Ammonium dihydrogen phosphate 7758-94-3, Ferrous chloride 7783-28-0, Diammonium hydrogen phosphate 10045-86-0, Ferric phosphate 10124-31-9, Ammonium phosphate 10377-48-7, Lithium sulfate 10377-52-3, Lithium phosphate 13453-80-0, Lithium dihydrogen phosphate 14940-41-1, Ferrous phosphate 17375-37-0, Manganese 18130-42-2, Cobalt oxalate 33943-39-4, carbonate Dilithium hydrogen phosphate 52767-99-4, Ammonium iron phosphate
- RL: PEP (Physical, engineering or chemical process); PROC (Process) (method for preparing spherical or quasi-spherical metal lithium phosphate)
- IT 13826-59-0P, Lithium manganese phosphate 15365-14-7P
  - , Ferrous lithium phosphate 153456-60-1P
    - 554453-38-2P, Iron lithium manganese phosphate
    - 915945-24-3P, Copper iron lithium zinc phosphate
    - 915945-25-4P, Cobalt iron lithium manganese phosphate
    - RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)
      - (method for preparing spherical or quasi-spherical metal lithium phosphate)
- L20 ANSWER 10 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN AN 2005:589401 HCAPLUS Full-text

- DN 143:118019
- ΤI Process for preparing electroactive insertion compounds and electrode materials obtained therefrom
- Gauthier, Laurent; Gauthier, Michel; Lavoie, Donald; Michot, IN Christophe; Ravet, Nathalie
- PA Universite De Montreal, Can.; Centre National de la Recherche
- SO
- DT
- LA

SO DT	Scientifique; Phostech Lithium Inc.  PCT Int. Appl., 50 pp.  CODEN: PIXXD2										пе						
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		ENT I	NO.			KIN	D	DATE			APPI.	TCAT	ION :	NO.		D	ATE
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	CA	2550	496			A1		2005	0707		CA 2	004-	2550	496			
																2	00412
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	EP	1702	373			A1		2006	0920		EP 2	004-	8023	57			
																2	00412
																2	2
		R:	ΑT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	ΙT,	LI,	LU,	NL,	SE,	MC,
				IE,	SI,			RO,							PL,	SK,	IS
	CN	1926	701			A		2007	0307		CN 2	004-	8004	1561			
																	00412
																2	2
	JP	2007	5157	62		T		2007	0614		JP 2	006-	5458	70			
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																2	2
	US	2006	0127	767		A1		2006	0615		US 2	005-	5364	31			
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KR 2007019972 A 20070216 KR 2006-714689

200607 21

PRAI US 2003-531606P P 20031223 WO 2004-CA2182 W 20041222

AB The invention relates to a process for preparing an at least partially lithiated transition metal oxyanion-based lithium-ion reversible electrode material, which comprises providing a precursor of the lithium-ion reversible electrode material, heating the precursor, melting same at a temperature sufficient to produce a melt comprising an oxyanion containing liquid phase, cooling the melt under conditions to induce solidification thereof and obtain a solid electrode that is capable of reversible lithium ion deinsertion/insertion cycles for use in a lithium battery. The invention also relates to lithiated or partially lithiated oxyanion-based-lithium-ion reversible electrode materials obtained by the aforesaid process.

IT 13816-45-0, Triphylite

RL: DEV (Device component use); USES (Uses)

(process for preparing electroactive insertion compds. and electrode  $% \left( 1\right) =\left( 1\right) \left( 1\right$ 

materials obtained therefrom)

RN 13816-45-0 HCAPLUS

CN Triphylite (FeLi(PO4)) (7CI, 9CI) (CA INDEX NAME)

● Fe(II)

■ T.i

IT 554453-38-2P, Iron lithium manganese phosphate

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

(process for preparing electroactive insertion compds. and electrode  $% \left( 1\right) =\left( 1\right) \left( 1\right$ 

materials obtained therefrom)

RN 554453-38-2 HCAPLUS

CN Phosphoric acid, iron lithium manganese(2+) salt (9CI) (CA INDEX NAME)

- Te(x)
  - ●x Li
- Mn(II)
- IT 33943-39-4

RL: RCT (Reactant); RACT (Reactant or reagent)

(process for preparing electroactive insertion compds. and electrode  $% \left( 1\right) =\left( 1\right) \left( 1\right$ 

materials obtained therefrom)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

♠ 2 T.i.

IT 13826-59-0P, Lithium manganese phosphate 15365-14-7DP, chromium- and molybdenum-doped 19414-36-9P, Iron lithium manganese phosphate ((Fe,Mn)Li(PO4)) 643752-34-5P, Iron lithium magnesium phosphate (Fe0.95LiMg0.05(PO4))

RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(process for preparing electroactive insertion compds. and electrode  $% \left( 1\right) =\left\{ 1\right\} =\left\{ 1\right$ 

materials obtained therefrom)

RN 13826-59-0 HCAPLUS

CN Phosphoric acid, lithium manganese(2+) salt (1:1:1) (9CI) (CA INDEX NAME)

● Li

Mn(II)

RN 15365-14-7 HCAPLUS CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

CN Iron lithium manganese phosphate ((Fe,Mn)Li(PO4)) (CA INDEX NAME)

Component		Ratio	- 1	Component
			- 1	Registry Number
	=+=		===+=	
04P	- 1	1	- 1	14265-44-2
Mn	- 1	0 - 1	- 1	7439-96-5
Li	- 1	1	- 1	7439-93-2
Fe	- 1	0 - 1	- 1	7439-89-6

RN 643752-34-5 HCAPLUS

CN Iron lithium magnesium phosphate (Fe0.95LiMg0.05(PO4)) (CA INDEX NAME)

Component	1	Ratio	Component Registry Number
	=+=		
04P		1	14265-44-2
Mg		0.05	7439-95-4
Li	- 1	1	7439-93-2
Fe	- 1	0.95	7439-89-6

IT 15365-14-7

RL: TEM (Technical or engineered material use); USES (Uses) (process for preparing electroactive insertion compds. and electrode

materials obtained therefrom)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

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IC ICM H01M004-04
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ICS C25B011-04; C07F001-02; H01M004-48; C01B025-26

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 49

IT 13816-45-0, Triphylite

RL: DEV (Device component use); USES (Uses)

(process for preparing electroactive insertion compds. and electrode

materials obtained therefrom)

IT 554453-38-2P, Iron lithium manganese phosphate

RL: DEV (Device component use); SPN (Synthetic preparation); PREP (Preparation); USES (Uses)

 $\begin{tabular}{ll} \hline \end{tabular}$  (process for preparing electroactive insertion compds. and electrode

materials obtained therefrom)

554-13-2, Lithium carbonate 1308-38-9, Chromium trioxide (Cr203), ΤТ reactions 1309-37-1, Ferric oxide, reactions 1310-65-2, Lithium hydroxide 1313-13-9, Manganese dioxide, reactions 1313-27-5, Molybdenum trioxide, reactions 1314-56-3, Phosphorus pentoxide, reactions 1317-61-9, Iron oxide (Fe304), reactions 1345-25-1, Ferrous oxide, reactions 7439-89-6, Iron, reactions 7447-41-8. Lithium chloride, reactions 7722-76-1, Ammonium dihydrogen phosphate 7783-28-0, Ammonium hydrogen phosphate 7789-24-4. Lithium fluoride, reactions 10045-86-0, Ferric phosphate 10377-52-3, Lithium phosphate 13453-80-0, Lithium dihydrogen phosphate 14940-41-1, Ferrous phosphate 33943-39-4 RL: RCT (Reactant); RACT (Reactant or reagent)

(process for preparing electroactive insertion compds. and

materials obtained therefrom)

IT 13826-59-0P, Lithium manganese phosphate
15365-14-7DP, chromium- and molybdenum-doped

19414-36-9P, Iron lithium manganese phosphate

((Fe,Mn)Li(PO4)) 643752-34-5P, Iron lithium magnesium phosphate (Fe0.95LiMg0.05(PO4))

RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(process for preparing electroactive insertion compds. and electrode

materials obtained therefrom)

IT 15365-14-7

RL: TEM (Technical or engineered material use); USES (Uses) (process for preparing electroactive insertion compds. and electrode

materials obtained therefrom)

RE.CNT 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

L20 ANSWER 11 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2004:492320 HCAPLUS Full-text

DN 141:26150

TI Preparation of a cathode material for secondary batteries

IN Franger, Sylvain; Martinet, Sebastien; Le Cras, Frederic; Bourbon, Carole

PA Commissariat A L'energie Atomique, Fr.

SO Fr. Demande, 33 pp.

CODEN: FRXXBL

DT Patent LA French

FAN.CNT 1

FAN.		TENT I	NO.			KIN	D -	DATE			APE	PLI	CAT	ION	NO.		D	ATE
ΡI	FR	2848	- 549			A1		2004	0618		FR	20	02-	1591	5			00212
		2848									To To	20	ns .	ED E O	170		1	6
	WO	2004056702				A2 20040708				WO 2003-FR501/2					_	00312		
	WO	2004 W:				А3		2004	0819								-	
			IE,	IT,	LU,	MC,	NL,	PT,	DE, RO,	SE,	SI	į, :	SK,	TR	·	GB,	GR,	HU,
	EP	1572	585			A2		2005	0914		EP	20	03-	8099	85		_	00312
			PT,	IE,	SI,	FI,	RO,	CY,	TR,	BG,	CZ	2, 1	EE,	HU,	SK	NL,	SE,	MC,
	CN	1726	167			A		2006	0125		CN	20	03-	8010	6132			00312
	CN	1003	7647	4		С		2008	0326									
	JP	2006	5114	21		T		2006	0406		JP	20	04-	5615	77		_	00312
	US	2006	0204	848		A1		2006	0914		US	20	06-	5379	47		_	9
																		00602 6
PRAI		2002																
	WO	2003	-FR5	0172		W		2003	1215									

AB A cathode material for secondary batteries is prepared having the general formula AMXO4 with A being an alkali metal, especially Li or Na, M being a transition metal, especially trivalent Mn, Fe, Ni, or Co, and X being Si, S, Al, Ge, As, Mo, preferably P. The material is prepared by reacting a complex of M bound to an organic ligand, such as nitrilotriacetic acid or EGTA, with a metal salt, especially Li2HPO4. The anode of the secondary battery is made of Li4Ti5012. 15365-14-7P, Iron lithium phosphate felipo4
RL: CPS (Chemical process); DEV (Device component use); PEP

(Physical, engineering or chemical process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process); USES (Uses) (cathode material; preparation of cathode material for secondary batteries)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

IΤ

● Fe(II)

● Li

IT 33943-39-4, Lithium phosphate (Li2HPO4)

RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent) (preparation of cathode material for secondary batteries)

RN 33943-39-4 HCAPLUS

CN Phosphoric acid, dilithium salt (8CI, 9CI) (CA INDEX NAME)

■ 2 T.-i

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IC ICM C01D001-02
ICS H01M004-48; H01M004-24; H01M010-24; C01D015-02; G02F001-15
CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
Section cross-reference(s): 49
IT 15365-14-7P, Iron lithium phosphate felipo4
RL: CPS (Chemical process); DEV (Device component use); PEP
(Physical, engineering or chemical process); SPN (Synthetic
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- RL: CPS (Chemical process); DEV (Device component use); PEP (Physical, engineering or chemical process); SPN (Synthetic preparation); PREP (Preparation); PROC (Process); USES (Uses) (cathode material; preparation of cathode material for secondary batteries)

  IT 67-42-5. EGTA 139-13-9, Glycine, N.N-bis(carboxymethyl)-
- 10028-22-5, Iron sulfate fe2(S04)3 33943-39-4, Lithium phosphate (Li2HPO4)
  RL: CPS (Chemical process); PEP (Physical, engineering or chemical process); RCT (Reactant); PROC (Process); RACT (Reactant or reagent) (preparation of cathode material for secondary batteries)
- L20 ANSWER 12 OF 12 HCAPLUS COPYRIGHT 2009 ACS on STN
- AN 1997:295088 HCAPLUS Full-text
- DN 127:43969
- OREF 127:8214h,8215a
- TI Interactions in the M2O-P2O5-NiO system (M = Li, Na, K)
- AU Nagornyi, P. G.; Petrenko, O. V.; Slobodyanik, N. S.
- CS Nats. Univ. im. Tarasa Shevchenka, Kiev, Ukraine
- SO Ukrainskii Khimicheskii Zhurnal (Russian Edition) (1996), 62(11-12), 14-18
  - CODEN: UKZHAU; ISSN: 0041-6045
- PB Institut Obshchei i Neorganicheskoi Khimii NAN Ukrainy
- DT Journal
- LA Russian
- AB The reactions of NiO with melts of MH2PO4 or M2HPO4 (M = Li, Na, K) were studied by the isothermal saturation and slow cooling methods at  $1000-750^{\circ}$ . The composition of the products was determined and the products were characterized by x-ray phase anal., IR spectra and derivatog. anal.
- IT 13977-83-8P, Lithium nickel phosphate (LiNiPO4)
  RL: SPN (Synthetic preparation); PREP (Preparation)
  (preparation by reaction of nickel oxide with alkali metal phosphate
- melts)
- RN 13977-83-8 HCAPLUS

10/537 947

■ 1. i

Ni(II)

■2 Lif

78-9 (Inorganic Chemicals and Reactions) CC ΙT 13977-83-8P, Lithium nickel phosphate (LiNiPO4) 14448-18-1P, Nickel phosphate (Ni2P2O7) 40437-73-8P, Nickel sodium phosphate (NiNa(PO3)3) 63090-58-4P, Nickel ultraphosphate 68877-75-8P, Nickel potassium metaphosphate oxide (NiP4011) (Ni4K2(PO3)4O3) 157098-42-5P, Nickel sodium diphosphate (NiNaP2O7) 190662-02-3P, Nickel sodium metaphosphate oxide (Ni2Na3(PO3)302) 190662-04-5P, Nickel sodium metaphosphate oxide (Ni3Na4(PO3)4O3) 190662-05-6P, Nickel potassium metaphosphate oxide (Ni3K6(PO3)802) 190662-07-8P, Nickel potassium metaphosphate oxide (Ni4K4(PO3)603) 190662-10-3P, Lithium nickel metaphosphate oxide (Li4Ni5(PO3)1002) 190662-12-5P, Lithium nickel metaphosphate oxide (Li2Ni3(PO3)60) RL: SPN (Synthetic preparation); PREP (Preparation) (preparation by reaction of nickel oxide with alkali metal

## phosphate

melts)

1313-99-1, Nickel oxide (NiO), reactions 7558-79-4, Disodium phosphate 7558-80-7, Sodium dihydrogen phosphate 7778-77-0, Potassium dihydrogen phosphate 13453-80-0, Lithium dihydrogen phosphate 25681-80-5, Dipotassium, reactions 33943-39-4, Dilithium phosphate

RL: RCT (Reactant); RACT (Reactant or reagent) (reaction of nickel oxide with alkali metal phosphate melts)

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- L30 ANSWER 1 OF 5 HCAPLUS COPYRIGHT 2009 ACS on STN
- AN 2008:843351 HCAPLUS Full-text
- DN 149:227142
- TΙ Method for synthesizing LixMy(PO4)z compounds under electron beam irradiation
- IN Zhao, Bing; Jiao, Zheng; Wu, Minghong; Yan, Jing; Zhong, Mingyang; He, Yagin; Jiang, Yong; Sun, Yufei; Wang, Song
- Shanghai University, Peop. Rep. China PA
- Faming Zhuanli Shenging Gongkai Shuomingshu, 6pp. SO CODEN: CNXXEV
- DT Patent
- Chinese T.A

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	CN 101214942	A	20080709	CN 2008-10032410	200801

PRAI CN 2008-10032410

20080108

The title compds. have a formula of LixMy(PO4)z, wherein M is one or AB two of Fe, Co, Ni, Mn, V, Cu, Ti, Cr, Mg and Zn. The compds. are synthesized by the following steps of: (1) weighing soluble M salt and phosphorus-containing compound, dissolving in deionized water, adding proper complexing agent, and then adding soluble Li salt under stirring, (2) adding suitable dilute base solution to adjust pH to 6.5-7, and ultrasonic-vibrating for 5-10 min, (3) electron beamirradiating at 20-40 Mrad in an electron accelerator (power 2.5 MeV and current 40 mA), (4) washing, centrifugating, and repeating many times to remove unreacted ion and complexing agent, (5) vacuumdrving, and (6) thermally treating in a tubular furnace at 400-600° for 5-10 h, and naturally cooling to obtain the final product with particle size of 50-100 nm. The concentration ratio of complexing agent to M ion is (0.1-1):1. The M salt is M nitrate or sulfate. The P-containing compound is phosphoric acid, diammonium hydrogen phosphate or ammonium dihydrogen phosphate. The Li salt is lithium hydroxide, lithium chloride, lithium sulfate or lithium carbonate. The complexing agent is disodium ethylenediaminetetraacetate, citric acid or aminotriacetic acid. The product can be used to prepare cathode materials of lithium ion batteries.

IΤ 139-13-9

RL: NUU (Other use, unclassified); USES (Uses)

(method for synthesizing LixMy(PO4)z compds. under electron beam irradiation)

RN 139-13-9 HCAPLUS

Glycine, N,N-bis(carboxymethyl) - (CA INDEX NAME) CN

IT 15365-14-7P, Iron lithium phosphate (FeLiPO4)

478819-84-0P, Iron lithium magnesium phosphate (FeLi0.99Mq0.01(PO4))

RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or

engineered material use); PREP (Preparation); USES (Uses) (method for synthesizing LixMy(PO4)z compds. under electron beam

irradiation)

RN 15365-14-7 HCAPLUS

Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX CN NAME)

- Fe(II)
  - Li

RN 478819-84-0 HCAPLUS

CN Iron lithium magnesium phosphate (FeLi0.99Mg0.01(PO4)) (CA INDEX NAME)

Component	I I	Ratio	l I R	Component egistry Number
	==+==		===+===	
04P	1	1	1	14265-44-2
Mg	1	0.01	1	7439-95-4
Li	1	0.99	1	7439-93-2
Fe	- 1	1	1	7439-89-6

- CC 49-3 (Industrial Inorganic Chemicals)
   Section cross-reference(s): 52
- IT 64-17-5, Ethanol, uses 77-92-9, Citric acid, uses 139-13-9 139-33-3, Disodium EDTA 1310-58-3, Potassium hydroxide, uses RL: NUU (Other use, unclassified); USES (Uses)

(method for synthesizing LixMy(PO4)z compds. under electron beam irradiation)

- IT 15365-14-7P, Iron lithium phosphate (FeLiPO4) 84159-18-2P, Lithium vanadium phosphate (Li3V2(FO4)3) 478819-84-0P, Iron lithium magnesium phosphate (FeLi0.99Mg0.01(FO4))
  - RL: PRP (Properties); SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)

(method for synthesizing LixMy(PO4)z compds. under electron beam irradiation)

- L30 ANSWER 2 OF 5 HCAPLUS COPYRIGHT 2009 ACS on STN
- AN 2005:368529 HCAPLUS Full-text
- DN 142:433067
- TI Manufacture of powdered anode active mass, the powdered electrode active mass, the electrode, and lithium battery

IN Saito, Mitsumasa; Toge, Yoshivuki

PA Sumitomo Osaka Cement Co., Ltd., Japan

SO Jpn. Kokai Tokkyo Koho, 15 pp.

CODEN: JKXXAF
DT Patent

7 Facenc

LA Japanese

r AN.	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PΙ	JP 2005116393	A	20050428	JP 2003-350632	

20031009

200310

PRAI JP 2003-350632

AB The powdered anode active mass is LixMyM'zPO4 (M = Fe, Co, Mn, Ni, Cr, and/or Cu; M' = Mg, Ca, Ba, Ti, Zn, b, Al, Ga, In, Si, Ge, Sc, Y, and/or rare earth metal), and is prepared by spraying a solution, dispersion, or suspension containing LiOH, sources of M and M', H3PO4 and/or phosphate salt, reaction inhibitor for LiOH and H3PO4 and/or phosphate, and reaction inhibitor for M and M' sources and H3PO4 and/or phosphate in a high temperature atmospheric to obtain a precursor, and firing the precursor.

IT 13824-63-OP, Cobalt lithium phosphate (CoLiPO4) 13826-59-OP, Lithium manganese phosphate (LiMnPO4) 15365-14-7P, Iron lithium phosphate (FeLiPO4)

RL: DEV (Device component use); IMF (Industrial manufacture); PREP (Preparation); USES (Uses)

(compns. and manufacture of powdered anode active mass for secondary

lithium batteries)

RN 13824-63-0 HCAPLUS

CN Phosphoric acid, cobalt(2+) lithium salt (8CI, 9CI) (CA INDEX NAME)

Oc(II)

● Li

RN 13826-59-0 HCAPLUS

CN Phosphoric acid, lithium manganese(2+) salt (1:1:1) (9CI) (CA INDEX NAME)

● Li

Mn(II)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

IT 139-13-9, Nitrilotriacetic acid

RL: NUU (Other use, unclassified); USES (Uses)

(in manufacture of powdered anode active mass by high temperature mist spraying  $% \left( 1\right) =\left( 1\right) +\left(  

and firing for secondary lithium batteries)

```
RN
    139-13-9 HCAPLUS
CN Glycine, N.N-bis(carboxymethyl) - (CA INDEX NAME)
          CH2-CO2H
 HO2C-CH2-N-CH2-CO2H
IC ICM H01M004-58
    ICS C01B025-45; H01M004-02; H01M010-40
    52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
    13824-63-0P, Cobalt lithium phosphate (CoLiPO4)
IΤ
    13826-59-0P, Lithium manganese phosphate (LiMnPO4)
     15365-14-7P, Iron lithium phosphate (FeLiPO4)
    RL: DEV (Device component use); IMF (Industrial manufacture); PREP
     (Preparation); USES (Uses)
        (compns. and manufacture of powdered anode active mass for
secondary
       lithium batteries)
ΙT
    77-92-9, Citric acid, uses 87-69-4, Tartaric acid, uses
    139-13-9, Nitrilotriacetic acid
     RL: NUU (Other use, unclassified); USES (Uses)
        (in manufacture of powdered anode active mass by high temperature
mist spraying
       and firing for secondary lithium batteries)
L30 ANSWER 3 OF 5 HCAPLUS COPYRIGHT 2009 ACS on STN
AN
    2003:719230 HCAPLUS Full-text
DN
    139:241693
TT
    Selective herbicide compositions comprising transition metal
    chelates
    Sedun, Frederick S.; Taylor, Kim F.; Wilson, Cameron D.; Parker,
IN
    Diana L.; Almond, David S.
PA W. Neudorff G.m.b.H. K.-G., Germany
SO PCT Int. Appl., 33 pp.
    CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 2
    PATENT NO.
                KIND DATE APPLICATION NO. DATE
PI WO 2003073856 A1 20030912 WO 2003-EP2069
```

AB

IΤ

RN

CN

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GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ,
            LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
            NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ,
             TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
            BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK,
            EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI,
            SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE,
             SN, TD, TG
                               20030912
                                         CA 2003-2477493
    CA 2477493
                         A1
                                                                  200302
                                                                  28
    AII 2003211724
                        Δ1
                              20030916
                                          AII 2003-211724
                                                                  200302
                                                                  28
    EP 1489909
                        A1
                               20041229 EP 2003-743354
                                                                  200302
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC,
            PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU,
             SK
PRAI US 2002-361217P
                               20020301
                        P
    WO 2003-EP2069
                        W
                               20030228
     An environmentally safe selective herbicide is provided that includes
     at least one metal component and at least one chelating agent. The
     metal component can have a variety of forms, but is preferably in the
     form of a metal salt, a metal chelate, or combinations thereof. The
     chelating agent can also have a variety of forms, but is preferably
     in the form of a metal chelate, a salt, an acid, or combinations
     thereof.
    67-42-5 139-13-9, Nitrilotriacetic acid
    RL: AGR (Agricultural use); BSU (Biological study, unclassified);
    BIOL (Biological study); USES (Uses)
        (chelating agent in selective herbicide compns. comprising
```

6,9-Dioxa-3,12-diazatetradecanedioic acid, 3,12-bis(carboxymethyl)-

AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD,

28

$$\begin{array}{c} {\rm CH_{2}-Co_{2}H} & {\rm CH_{2}-Co_{2}H} \\ {\rm Ho_{2}C-CH_{2}-N-CH_{2}-CH_{2}-O-CH_{2}-CH_{2}-O-CH_{2}-CH_{2}-N-CH_{2}-Co_{2}H} \end{array}$$

transition metal chelates)

67-42-5 HCAPLUS

(CA INDEX NAME)

RN 139-13-9 HCAPLUS CN Glycine, N,N-bis(carboxymethyl)- (CA INDEX NAME)

CH2-CO2H HO2C-CH2-N-CH2-CO2H

IT 596113-31-4

RL: AGR (Agricultural use); BSU (Biological study, unclassified); BIOL (Biological study); USES (Uses)

(selective herbicide compns. comprising)

RN 596113-31-4 HCAPLUS

Glycine, N,N'-1,2-ethanediylbis[N-(carboxymethyl)-, tetrasodium salt, mixt. with iron(2+) sulfate (1:1) (9CI) (CA INDEX NAME)

CM 1

CN

CRN 7720-78-7 CMF Fe . H2 O4 S

но— 3— он

● Fe(II)

CM 2

CRN 64-02-8

CMF C10 H16 N2 O8 . 4 Na

4 Na

IC ICM A01N059-16

ICS A01N059-20; A01N037-44; A01N037-32; A01N043-58

CC 5-3 (Agrochemical Bioregulators) IΤ

56-40-6, Glycine, biological studies 56-41-7, L-Alanine, biological studies 56-45-1, L-Serine, biological studies 56-84-8. Aspartic acid, biological studies 56-85-9, L-Glutamine, biological studies 56-86-0, Glutamic acid, biological studies 56-87-1, L-Lysine, biological studies 60-00-4, Ethylenediaminetetraacetic acid, biological studies 60-18-4, L-Tyrosine, biological studies 61-90-5, L-Leucine, biological studies 63-68-3, L-Methionine, biological studies 64-02-8, Glycine, N, N'-1, 2-ethanediylbis[N-(carboxymethyl)-, tetrasodium salt 64-19-7, Acetic acid, biological studies 67-42-5 67-43-6, Diethylenetriaminepentaacetic acid 69-72-7, Salicylic acid, biological studies 70-47-3, L-Asparagine, biological studies 71-00-1, L-Histidine, biological studies 72-18-4, L-Valine, biological studies 72-19-5, L-Threonine, biological studies 73-32-5, L-Isoleucine, biological studies 74-79-3, L-Arginine, biological studies 77-92-9, Citric acid, biological studies 80-69-3, Tartronic acid 87-69-4, Tartaric acid, biological studies 93-62-9 97-65-4, Itaconic acid, biological studies 102-71-6, Triethanolamine, biological studies 107-15-3, Ethylenediamine, biological studies 110-15-6, Succinic acid, biological studies 110-44-1, Sorbic acid 110-94-1, Glutaric acid 110-99-6, Diglycolic acid 111-42-2, Diethanolamine, biological studies 112-24-3 139-13-9, Nitrilotriacetic acid 141-43-5, Monoethanolamine, biological studies 142-73-4, Iminodiacetic acid 147-85-3, L-Proline, biological studies 149-91-7, Gallic acid, biological studies 150-25-4, Diethanol glycine 482-54-2 498-23-7, Citraconic acid 499-12-7, Aconitic acid 499-83-2, Dipicolinic acid 528-94-9, Ammonium salicylate 541-50-4, Diacetic acid, biological studies 688-57-3D, Ethylenediamine triacetic acid. alkovl derivs. 817-11-8. Nitrilotripropionic acid 1170-02-1 2809-21-4 3148-72-9 4408-64-4, Methyliminodiacetic acid 4408-81-5, 1,2-Diaminopropanetetraacetic acid 6419-19-8, Aminotri (methylenephosphonic acid) 7408-20-0, Iminodisuccinic acid 13073-35-3, Ethionine 13288-40-9 13598-36-2, Phosphonic

TΤ

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20846-91-7 23351-51-1, Glucoheptonic acid 29578-05-0,
Methyl glycine diacetic acid 32013-58-4 33872-70-7.
Hydroxyethylenediaminetriacetic acid 40623-42-5 58976-65-1
87339-38-6, Hydroxyiminodiacetic acid 89198-07-2 133677-59-5
148124-42-9
            167613-87-8
RL: AGR (Agricultural use); BSU (Biological study, unclassified);
BIOL (Biological study); USES (Uses)
   (chelating agent in selective herbicide compns. comprising
   transition metal chelates)
7439-89-6D, Iron, chelates 7440-02-0D, Nickel, chelates
7440-50-8D, Copper, chelates
                            7440-66-6D, Zinc, chelates
15009-37-7 17084-02-5 17099-81-9
                                     51395-10-9, Copper EDTA
55448-21-0, ZnEDTA 167256-48-6 167256-48-6D, mixture with beet
molasses 187165-07-7 596113-16-5 596113-17-6 596113-18-7
596113-19-8
            596113-20-1
                         596113-21-2
                                       596113-22-3
                                                    596113-23-4
596113-24-5 596113-25-6
                         596113-26-7
                                       596113-27-8
                                                    596113-28-9
596113-29-0 596113-30-3 596113-31-4 596113-32-5
596113-33-6 596113-34-7 596113-35-8 596113-36-9 596113-37-0
596113-38-1 596113-39-2 596113-40-5 596113-42-7
                                                    596113-43-8
596113-44-9 596113-45-0
RL: AGR (Agricultural use); BSU (Biological study, unclassified);
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BIOL (Biological study); USES (Uses) (selective herbicide compns. comprising)

RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L30 ANSWER 4 OF 5 HCAPLUS COPYRIGHT 2009 ACS on STN
- AN 2003:437426 HCAPLUS Full-text
- DN 139:278928
- TI Comparison between different LiFePO4 synthesis routes and their influence on its physico-chemical properties
- AU Franger, Sylvain; Le Cras, Frederic; Bourbon, Carole; Rouault, Helene
- CS DRT/DTEN/SCSE/LSEM, Commissariat a l'Energie Atomique, Grenoble, 38054, Fr.
- SO Journal of Power Sources (2003), 119-121, 252-257 CODEN: JPSODZ: ISSN: 0378-7753
- PB Elsevier Science B.V.
- DT Journal
- LA English
- AB LiFePO4 powders were synthesized using solid state reactions at high temps., co-precipitation in aqueous medium, hydrothermal synthesis or mechanochem. activation. The samples were characterized by XRD, chemical titration and their electrochem. performance were studied for cycling behavior. It is advantageous to introduce an electronic conductor precursor (typically a sucrose) during or after the

synthesis to overcome the poor charge transfer associated with LiFePO4.

1T 139-13-9D, Nitrilotriacetic acid, iron complexes
RL: CPS (Chemical process); PEP (Physical, engineering or chemical
process); PROC (Process)

(in synthesis of LiFePO4 for cathodes of lithium batteries)

RN 139-13-9 HCAPLUS

CN Glycine, N, N-bis(carboxymethyl) - (CA INDEX NAME)

IT 15365-14-7P, Iron lithium phosphate (FeLiPO4)
RL: DEV (Device component use); PNU (Preparation, unclassified); PRP
(Properties); PREP (Preparation); USES (Uses)

(influence of synthesis route on properties of LiFePO4 cathode material for lithium batteries)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 72
- IT 139-13-9D, Nitrilotriacetic acid, iron complexes
  7439-89-6D, Iron, nitrilotriacetic acid complexes
  RL: CPS (Chemical process); PEP (Physical, engineering or chemical

ΙT

process); PROC (Process)

(in synthesis of LiFePO4 for cathodes of lithium batteries) 15365-14-7P, Iron lithium phosphate (FeLiPO4)

RL: DEV (Device component use); PNU (Preparation, unclassified); PRP (Properties); PREP (Preparation); USES (Uses)

(influence of synthesis route on properties of LiFePO4 cathode material for lithium batteries)

RE.CNT 10 THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L30 ANSWER 5 OF 5 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 1998:127054 HCAPLUS Full-text

DN 128:148754

OREF 128:29109a,29112a

- TI Synthesis and Crystal Structure of Maricite and Sodium Iron(III) Hydroxyphosphate
- AU Bridson, John. N.; Quinlan, Sean; Tremaine, Peter R.
- CS Department of Chemistry, Memorial University of Newfoundland, St. John's, NF, A1B 3X7, Can.
- SO Chemistry of Materials (1998), 10(3), 763-768 CODEN: CMATEX; ISSN: 0897-4756
- PB American Chemical Society
- DT Journal
- LA English
- AB Maricite (NaFePO4) and sodium iron hydroxyphosphate (SIHP) are recently discovered iron(II) and iron(III) compds. that play a major role in phosphate hideout and corrosion in high-pressure boilers. This paper reports a novel method for synthesizing maricite by thermally decomposing the complex of aqueous iron(III) nitrilotriacetic acid at 200° and methods for the hydrothermal synthesis of SIHP from Fe304 or FeP04. The crystal structure of maricite is identical to the impure natural mineral. The x-ray diffraction pattern of SIHP is consistent with an orthorhombic unit cell containing 8 units Na3Fe(PO4)2  $\cdot$  (Na2(1-x)H2xO), with x = 0.226  $\pm$ 0.025. This is similar, but not identical, to Na4Fe(OH)(PO4)2·1/3NaOH proposed by Ziemniak and Opalka. The main structural feature is a chain of iron(III) ions linked by bridging oxygens and phosphate bridges. The iron phosphate chains are held together by sodium ions in the ratio 3Na/Fe/2(PO4). Charge balance with the O2- bridge between each iron ion is maintained by Na+ and H+ ions located in a relatively open cage in the phosphate lattice. The structure is significant in that it explains the variable stoichiometry observed in powder diffraction patterns, and it identifies the stoichiometry of H and O, which cannot be determined from solubility studies.
- IT 139-13-9, Nitrilotriacetic acid
  RL: RCT (Reactant); RACT (Reactant or reagent)

(for preparation of sodium iron phosphate (FeNaPO4)) RN 139-13-9 HCAPLUS

CN Glycine, N, N-bis(carboxymethyl) - (CA INDEX NAME)

IT 53602-70-3P, Iron sodium phosphate (FeNaPO4)
202334-18-7P, Iron sodium hydroxide oxide phosphate
(FeNa4.55(OH) 0.4500.55(PO4)2)
RL: PRP (Properties); SPN (Synthetic preparation); PREP

RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation)

(preparation and crystal structure)

RN 53602-70-3 HCAPLUS

CN Phosphoric acid, iron(2+) sodium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

Na

RN 202334-18-7 HCAPLUS

CN Iron sodium hydroxide oxide phosphate (FeNa4.55(OH)0.4500.55(PO4)2)
 (CA INDEX NAME)

Component	- 1	Ratio		Component
	- 1			Registry Number
	=+=		==+=	
0	- 1	0.55	- 1	17778-80-2
HO	- 1	0.45	- 1	14280-30-9
04P	- 1	2	- 1	14265-44-2

Na 4.55 7440-23-5 7439-89-6 Fe 1 CC 78-6 (Inorganic Chemicals and Reactions) Section cross-reference(s): 61, 75 ΙT 139-13-9. Nitrilotriacetic acid RL: RCT (Reactant); RACT (Reactant or reagent) (for preparation of sodium iron phosphate (FeNaPO4)) ΙT 53602-70-3P, Iron sodium phosphate (FeNaPO4) 202334-18-7P, Iron sodium hydroxide oxide phosphate (FeNa4.55(OH)0.4500.55(PO4)2) RL: PRP (Properties); SPN (Synthetic preparation); PREP (Preparation) (preparation and crystal structure) THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 27

ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d 135 1-9 bib abs hitstr hitind YOU HAVE REQUESTED DATA FROM FILE 'HCAPLUS' - CONTINUE? (Y)/N:y

L35 ANSWER 1 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2008:71588 HCAPLUS Full-text

DN 148:218494

TI High-safety lithium power battery and its assembly method

IN Liu, Lijun; Zhang, Baowen; Liu, Weiwei; Xu, Yan; Han, Lei; Zhou, Huansheng

PA Beijing China Powerel Battery Co., Ltd., Peop. Rep. China

SO Faming Zhuanli Shenqing Gongkai Shuomingshu, 16pp.

DT Patent

LA Chinese

FAN.CNT 1

AB

04

PRAI CN 2007-10146016

20070904

The battery has several unit cells, a shell storing the unit cells and elec. connecting the unit cells, and a flame retardant or heat-transfer medium filled in the shell. The assembly method is carried out by: (1) providing unit cells with required number, (2) placing

the unit cells in a shell, (3) elec. connecting the unit cells, (4) filling the flame retardant or heat-transfer medium into the shell. and (5) elec. connecting the unit cells to the shell, and sealing. The obtained battery has high safety.

126-73-8, Tributyl phosphate, uses 411234-54-3, ΙT

Iron Lithium phosphate

RL: TEM (Technical or engineered material use); USES (Uses) (structure and assembly method high-safety secondary lithium batteries)

126-73-8 HCAPLUS RN

Phosphoric acid tributyl ester (CA INDEX NAME) CN

RN 411234-54-3 HCAPLUS

CN Phosphoric acid, iron lithium salt (9CI) (CA INDEX NAME)

x Fe(x)

x Li

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) ΙT 108-78-1, Melamine, uses 115-86-6, Triphenyl phosphate 126-73-8, Tributyl phosphate, uses 383-63-1, Ethyl trifluoroacetate 1309-42-8, Magnesium hydroxide 7782-42-5, Graphite, uses 9003-53-6D, Polystyrene, Brominated 21645-51-2, Aluminum hydroxide, uses 39457-42-6, Lithium manganese oxide 411234-54-3, Iron Lithium phosphate

- RL: TEM (Technical or engineered material use); USES (Uses) (structure and assembly method high-safety secondary lithium batteries)
- L35 ANSWER 2 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN
- AN 2007:1329553 HCAPLUS Full-text
- DN 149:83426
- TI Synthesis and electrochemical properties of LiFePO4/C cathode material prepared by using tri-n-butyl phosphate as a multi-function agent
- AU Xu, Liang; Xu, Zhi-hui; Lai, Qiong-yu; Chen, Lian-mei; Ji, Xiao-yang CS College of Chemistry, Sichuan University, Chengdu, 610064, Peop.
- Rep. China SO Gongneng Cailiao (2007), 38(8), 1316-1319 CODEN: GOCAEA; ISSN: 1001-9731
- PB Gongneng Cailiao Bianjibu
- DT Journal
- LA Chinese
- AB LiFePO4/C cathode material had been synthesized by using tri-Bu phosphate (TBP) as a multi-function agent and polyethylene glycol 4000 (PEG-4000) as a surfactant and carbon source. The product structure, surface morphol. and chemical composition were detected by means of X-ray diffraction (XRD), scanning electron microscope (SEM) and X-ray photoelectron spectroscope (XPS). The results indicated that the sample prepared at 650 °C for 15 h was well crystallized with small and homogeneous global particles. In the voltage range of 2.5-4.3 V (vs Li), cyclic voltammetry tests showed there was a pair of sharp redox peaks with almost equal areas, indicating an excellent reversibility of the material. Electrochem. tests exhibited that the first discharge capacity was 158 mAh/g at the c.d. of 0.1 mA/ cm2. After 100th cycle, the specific discharge capacity was 153 mAh/g with only the loss of 3.3%. Even at the high c.d. of 4.0 mA/cm2, it still showed the discharge capacity close to 100 mAh/q.
- IT 126-73-8, Tri-n-butyl phosphate, uses
  - RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)
    - (synthesis and electrochem. properties of LiFePO4/C cathode material prepared by using tri-Bu phosphate as multi-function agent)
- RN 126-73-8 HCAPLUS
- CN Phosphoric acid tributyl ester (CA INDEX NAME)

10/537 947

IT 15365-14-7

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(synthesis and electrochem. properties of LiFePO4/C cathode material prepared by using tri-Bu phosphate as multi-function agent)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT 126-73-8, Tri-n-butyl phosphate, uses

RL: NUU (Other use, unclassified); PEP (Physical, engineering or chemical process); PROC (Process); USES (Uses)

(synthesis and electrochem. properties of LiFePO4/C cathode material prepared by using tri-Bu phosphate as multi-function agent)

IT 7440-44-0, Carbon, uses 15365-14-7 25322-68-3

RL: PRP (Properties); TEM (Technical or engineered material use); USES (Uses)

(synthesis and electrochem. properties of LiFePO4/C cathode material prepared by using tri-Bu phosphate as multi-function agent)

WO 2007-US63639 W

OS MARPAT 147:409729

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L35
    ANSWER 3 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN
AN
     2007:1089561 HCAPLUS Full-text
DN
    147:409729
TΙ
    Stabilized nonaqueous electrolytes for rechargeable batteries
    Xu, Wu; Deng, Zhongyi
IN
PΑ
    Ferro Corporation, USA
SO
    PCT Int. Appl., 22pp.
    CODEN: PIXXD2
DT
    Patent
LA
    English
FAN.CNT 1
    PATENT NO.
                       KIND DATE
                                     APPLICATION NO.
                                                                DATE
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PΤ
    WO 2007109435
                        A2 20070927
                                         WO 2007-US63639
                                                                 200703
                                                                 09
    WO 2007109435
                        A3 20080710
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA,
            CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
            GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE,
            KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA,
            MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG,
            PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY,
            TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
        RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU,
            IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK,
             TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN,
            TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG,
             ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA
    US 20070224515
                        A1 20070927 US 2006-387142
                                                                 200603
                                                                 22
    EP 1997183
                        A2 20081203 EP 2007-758213
                                                                 200703
                                                                 0.9
            AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU,
             IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI,
             SK, TR, AL, BA, HR, MK, RS
                             20081203 KR 2008-722785
     KR 2008105087
                        A
                                                                 200809
                                                                 18
PRAI US 2006-387142
                        A 20060322
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20070309

AB The invention relates to the use of aromatic phosphite compds. as stabilizers for nonaq. electrolytic solns. containing halogenated salts such as LiPF6 and LiBF4. The electrolyte containing such a phosphite exhibits excellent shelf life storage at ambient and high temps. The electrolytic solution is suitable for use in electrochem. cells such as lithium (ion) rechargeable batteries and

supercapacitors.

IT 126-73-8, Tributyl phosphate, uses 15365-14-7,
Iron lithium phosphate felipo4
RL: TEM (Technical or engineered material use); USES (Uses)
(stabilized nonaq. electrolytes for rechargeable batteries)
RN 126-73-8 HCAPLUS

CN Phosphoric acid tributyl ester (CA INDEX NAME)

$$n-BuO - P OBu-n$$
 $OBu-n$ 

RN 15365-14-7 HCAPLUS CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 76
- IT 57-57-8,  $\beta$ -Propiolactone 68-12-2, Dmf, uses 75-05-8, Acetonitrile, uses 78-40-0, Triethyl phosphate 79-20-9, Methyl

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acetate 96-47-9, 2-Methyltetrahydrofuran 96-48-0,
y-Butyrolactone 96-49-1, Ethylene carbonate 105-37-3,
Ethyl propionate 105-54-4, Ethyl butyrate 105-58-8, Diethyl
carbonate 105-66-8, Propyl butyrate 106-36-5, Propyl propionate
107-31-3, Methyl formate 108-29-2, 4-Methyl-Y-Butyrolactone
108-32-7, Propylene carbonate 109-21-7, Butyl butyrate 109-60-4,
Propyl acetate 109-94-4, Ethyl formate 109-99-9, Thf, uses
110-71-4, 1,2-Dimethoxyethane 110-74-7, Propyl formate 112-48-1,
1,2-Dibutoxyethane 115-86-6, Triphenyl phosphate 115-96-8,
Tris(2-chloroethyl)phosphate 123-86-4, Butyl acetate 123-91-1,
1,4-Dioxane, uses 126-73-8, Tributyl phosphate, uses
141-78-6, Ethyl acetate, uses 358-63-4,
Tris(2,2,2)trifluoroethyl)phosphate 512-56-1, Trimethyl phosphate
513-02-0, Triisopropyl phosphate 513-08-6, Tripropyl phosphate
542-28-9, δ-Valerolactone 542-52-9, Dibutyl carbonate
554-12-1, Methyl propionate 590-01-2, Butyl propionate 592-84-7,
Butyl formate 616-38-6, Dimethyl carbonate 623-42-7, Methyl
butyrate 623-53-0, Ethyl methyl carbonate 623-96-1, Dipropyl
carbonate 629-14-1, 1,2-Diethoxyethane 646-06-0, 1,3-Dioxolane
1301-96-8, Silver oxide (AgO) 1307-96-6, Cobalt oxide (CoO), uses
1309-60-0, Lead oxide (PbO2) 1309-64-4, Antimony oxide (Sb2O3),
uses 1310-53-8, Germanium oxide (GeO2), uses 1312-43-2, Indium
oxide (In203) 1313-99-1, Nickel oxide (NiO), uses 1314-13-2,
Zinc oxide (ZnO), uses 1314-27-8, Lead oxide (Pb2O3) 1314-41-6,
Lead oxide (Pb304) 1314-60-9, Antimony oxide (Sb205) 1317-36-8,
Lead oxide (PbO), uses 1330-78-5, Tritolyl phosphate 1332-81-6,
Antimony oxide (Sb2O4) 1345-25-1, Iron oxide (FeO), uses
1679-47-6, 2-Methyl-y-Butyrolactone 1679-49-8,
3-Methyl-y-Butyrolactone 2528-39-4, Trihexyl phosphate
4437-85-8, Butylene carbonate 7439-93-2, Lithium, uses
7440-44-0, Carbon, uses 7791-03-9, Lithium perchlorate
12002-97-0, Silver oxide (Ag203) 12022-46-7, Iron lithium oxide
(FeLiO2) 12030-22-7, Indium oxide (In2O) 12031-65-1, Lithium
nickel oxide (LiNiO2) 12031-95-7. Lithium titanium oxide
(Li4Ti5012) 12057-17-9, Lithium manganese oxide (LiMn204)
12057-30-6 12162-79-7, Lithium manganese oxide limno2
12190-79-3, Cobalt lithium oxide (CoLiO2) 12332-29-5, Iron lithium
nitride (FeLi3N2) 12338-02-2 12798-95-7 13843-81-7. Lithium
dichromate li2cr2o7 14024-11-4, Lithium tetrachloroaluminate
14283-07-9, Lithium tetrafluoroborate 14307-35-8, Lithium chromate
li2cro4 15365-14-7, Iron lithium phosphate felipo4
18282-10-5. Tin oxide (SnO2) 20619-16-3, Germanium oxide (GeO)
20667-12-3, Silver oxide (Ag20) 21324-40-3, Lithium
hexafluorophosphate 21651-19-4, Tin oxide (SnO) 25743-90-2
29935-35-1, Lithium hexafluoroarsenate 33454-82-9, Lithium
triflate 35363-40-7, Ethyl propyl carbonate 37186-88-2
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DT LA

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56525-42-9, Methyl propyl carbonate 61234-06-8, Lithium 80,
    silicon 20 atomic 61535-79-3, Lithium 20, tin 80 atomic
    62852-65-7 82906-17-0 97037-12-2 113443-18-8, Silicon oxide
    (SiO) 128975-24-6, Lithium manganese nickel oxide limn0.5ni0.5o2
    135573-53-4. Cobalt lithium nickel oxide co0-1lini0-1o2
    174421-80-8, Cobalt lithium nitride (Co0.4Li2.6N) 182442-95-1,
    Cobalt lithium manganese nickel oxide 184912-51-4, Copper lithium
    nitride (Cu0.4Li2.6N) 476300-71-7, Lithium carbide (LiC6)
    945544-42-3
    RL: TEM (Technical or engineered material use); USES (Uses)
       (stabilized nonag. electrolytes for rechargeable batteries)
    ANSWER 4 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN
    2006:1256691 HCAPLUS Full-text
    Nonaqueous electrolytic solution for electrochemical cells
    Xu, Wu; Deng, Zhongyi; Prabhu, Vaikunth S.; Bolomey, Pascal
   Ferro Corporation, USA
   PCT Int. Appl., 19pp.
    CODEN: PIXXD2
    Pat.ent.
    English
FAN.CNT 1
    PATENT NO. KIND DATE APPLICATION NO. DATE
    _____
                                         _____
                       A2 20061130 WO 2006-US15294
    WO 2006127192
                                                                200604
                                                                24
    WO 2006127192
                        A3 20071011
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA,
            CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI,
            GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM,
            KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG,
            MK. MN. MW. MX. MZ. NA. NG. NI. NO. NZ. OM. PG. PH. PL. PT.
            RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT,
            TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
        RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU,
            IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR,
            BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD,
            TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,
            ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA
    US 20060269845
                       A1 20061130 US 2005-138769
                                                                26
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- AB The invention relates to the use of an amine oxide as an additive in a nonaq. electrolytic solution The electrolytic solution is suitable for use in electrochem. cells such as lithium batteries and lithium ion batteries. Batteries using this electrolyte solution have long life and high capacity retention.
- RN 126-73-8 HCAPLUS
- CN Phosphoric acid tributyl ester (CA INDEX NAME)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

- Fe(II)
  - Li
- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
  IT 57-57-8, β-Propiolactone 68-12-2, Dmf, uses 75-05-8,
   Acetonitrile, uses 78-40-0, Triethyl phosphate 79-20-9, Methyl
   acetate 96-47-9, 2-Methyltetrahydrofuran 96-48-0,
   y-Butyrolactone 96-49-1, Ethylene carbonate 105-37-3,
   Ethyl propionate 105-54-4, Ethyl butyrate 105-58-8, Diethyl

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carbonate 105-66-8, Propyl butyrate 106-36-5, Propyl propionate
107-31-3, Methyl formate 108-29-2, 4-Methyl-γ-Butyrolactone
108-32-7, Propylene carbonate 109-21-7, Butyl butyrate 109-60-4,
Propyl acetate 109-94-4, Ethyl formate 109-99-9, Thf, uses
110-71-4, 1,2-Dimethoxyethane 110-74-7, Propyl formate 112-48-1,
1,2-DiButoxyethane 115-86-6, Triphenyl phosphate 115-96-8,
Tris(2-chloroethyl)phosphate 123-86-4, Butyl acetate 123-91-1,
1,4-Dioxane, uses 126-73-8, Tributylphosphate, uses
141-78-6, Ethyl acetate, uses 358-63-4,
Tris(2,2,2-trifluoroethyl)phosphate 512-56-1, Trimethyl phosphate
513-02-0, Triisopropyl phosphate 513-08-6, Tripropyl phosphate
542-28-9, \delta-Valerolactone 542-52-9, Dibutyl carbonate
554-12-1, Methyl propionate 590-01-2, Butyl propionate
Butyl formate 616-38-6, Dimethyl carbonate 623-42-7, Methyl
butyrate 623-53-0, Ethyl methyl carbonate 623-96-1, Dipropyl
carbonate 629-14-1, 1,2-Diethoxyethane 646-06-0, 1,3-Dioxolane
1330-78-5, Tritolyl phosphate 1679-47-6,
2-Methyl-γ-Butyrolactone
                         1679-49-8.
3-Methyl-y-Butyrolactone 2528-39-4, Trihexyl phosphate
4437-85-8, Butylene carbonate 7439-93-2, Lithium, uses
7440-44-0, Carbon, uses 7791-03-9, Lithium perchlorate
12022-46-7, Iron lithium oxide (FeLiO2) 12031-65-1, Lithium nickel
oxide (LiNiO2) 12031-95-7, Lithium titanium oxide (Li4Ti5O12)
12042-37-4, Alli 12057-17-9, Lithium manganese oxide (LiMn204)
12057-22-6, Lizn 12057-30-6 12162-79-7, Lithium manganese oxide
limno2 12190-79-3, Cobalt lithium oxide (CoLiO2) 12332-29-5,
Iron lithium nitride (FeLi3N2) 12338-02-2 13843-81-7, Dilithium
dichromate 14024-11-4, Lithium tetrachloroaluminate 14283-07-9,
Lithium tetrafluoroborate 14307-35-8, Lithium chromate
15365-14-7, Iron lithium phosphate felipo4 18424-17-4,
Lithium hexafluoroantimonate 21324-40-3, Lithium
hexafluorophosphate 25743-90-2 33454-82-9, Lithium triflate
35363-40-7, Ethyl propyl carbonate 56525-42-9, Methyl propyl
carbonate 61234-06-8, Lithium 80, silicon 20 atomic 62852-65-7,
Dilithium decachlorodecaborate(2) 82906-17-0 97037-11-1
97037-12-2 128975-24-6, Lithium manganese nickel oxide
limn0.5ni0.5o2 135573-53-4, Cobalt lithium nickel oxide
co0-1lini0-1o2 174421-80-8, Cobalt lithium nitride (Co0.4Li2.6N)
177997-11-4, Cobalt gallium lithium nickel oxide 177997-12-5,
Boron cobalt lithium nickel oxide 177997-13-6, Aluminum cobalt
lithium nickel oxide 180997-14-2. Cobalt lithium magnesium nickel
oxide 184912-51-4, Copper lithium nitride (Cu0.4Li2.6N)
244304-18-5, Cobalt lithium nickel silicon oxide 244304-20-9,
Cobalt lithium nickel titanium oxide
                                     244761-29-3, Lithium
bisoxalatoborate 291298-96-9 321201-33-6, Lithium
tris(oxalato)phosphate(1-) 346417-97-8, Cobalt lithium manganese
nickel oxide (Co0.33LiMn0.33Ni0.33O2) 383187-24-4 427879-42-3
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476300-71-7, Lithium carbide (LiC6) 913080-19-0 913080-20-3 916203-51-5, Lithium vanadium phosphate (Li3Y(PO4)) RL: TEM (Technical or engineered material use); USES (Uses) (nonaq, electrolytic solution for electrochem, cells)

L35 ANSWER 5 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN

AN 2006:1256638 HCAPLUS Full-text

DN 146:10718

TI Triazine compounds for removing acids and water from nonaqueous electrolytes for electrochemical cells

IN Deng, Zhongyi; Decker, Jerry L.; Xu, Wu; Sans, John R.; Bolomey, Pascal

PA Ferro Corporation, USA

SO U.S. Pat. Appl. Publ., 6pp.

CODEN: USXXCO

DT Patent

LA English

FAN.CNT 1

PΙ

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 20060269844	A1	20061130	US 2005-138907	
				200505
				26

PRAI US 2005-138907

20050526

OS MARPAT 146:10718

AB A process is provided to produce non-aqueous electrolytic solution for use in batteries having low acid content and low water content. The invention involves removing acids and water from non-aqueous electrolytic solns. typically found in lithium or lithium-ion batteries by using nitrogen-containing compds. such as triazines. After treatment by a triazine such as melamine, the concns. of acids and water in the electrolytic solns. are substantially decreased. The present invention provides a process to prepare extremely pure electrolytic solns. having low (<20 ppm) concns. of both water and acids.

II 126-73-8, Tributyl phosphate, uses 15365-14-7,

Iron lithium phosphate felipo4

RL: TEM (Technical or engineered material use); USES (Uses) (triazine compds. for removing acids and water from nonag.

electrolytes for electrochem. cells)

RN 126-73-8 HCAPLUS

CN Phosphoric acid tributyl ester (CA INDEX NAME)

RN 15365-14-7 HCAPLUS CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

• Fe(II)

Li

INCL 429325000; 429326000; 029623300; 029623200; 029623500 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 27 ΙT 57-57-8, Propiolactone 68-12-2, Dmf, uses 78-40-0, Triethvl phosphate 79-20-9, Methyl acetate 96-47-9, 2-Methvltetrahydrofuran 96-48-0, γ-Butyrolactone 96-49-1, Ethylene carbonate 105-37-3, Ethyl propionate, uses 105-54-4, Ethyl butyrate, uses 105-58-8, Diethyl carbonate 105-66-8, Propyl butyrate 106-36-5, Propyl propionate, uses 107-31-3, Methyl formate 108-29-2, 4-Methyl-v-Butyrolactone 108-32-7, Propylene carbonate 109-21-7, Butyl butyrate, uses 109-60-4, Propvl acetate 109-94-4, Ethvl formate 109-99-9, Thf. uses 110-71-4, 1,2-Dimethoxyethane 110-74-7, Propyl formate 112-48-1, 1,2-Dibutoxyethane 115-86-6, Triphenyl phosphate 115-96-8, Tris(chloroethyl) phosphate 123-86-4, Butyl acetate 123-91-1, 1,4-Dioxane, uses 126-73-8, Tributyl phosphate, uses 141-78-6, Ethyl acetate, uses 358-63-4,
Tris(2,2,2-trifluoroethyl) phosphate 512-56-1, Trimethyl phosphate 513-02-0, Triisopropyl phosphate 513-08-6, Tripropyl phosphate

542-28-9,  $\delta$ -Valerolactone 542-52-9, Dibutyl carbonate 554-12-1, Methyl propionate 590-01-2, Butyl propionate 592-84-7, Butyl formate 616-38-6, Dimethyl carbonate 623-42-7, Methyl butyrate 623-53-0, Ethyl methyl carbonate 623-96-1, Dipropyl carbonate 629-14-1, 1,2-Diethoxyethane 646-06-0, 1,3-Dioxolane 1330-78-5, Tritolyl phosphate 1679-47-6, 2-Methyl-γ-Butyrolactone 1679-49-8, 3-Methyl-y-Butyrolactone 2528-39-4, Trihexyl phosphate 4437-85-8, Butylene carbonate 7439-93-2, Lithium, uses 7439-93-2D, Lithium, salt 7440-44-0, Carbon, uses 7791-03-9, Lithium perchlorate 12022-46-7, Iron lithium oxide (FeLiO2) 12031-65-1, Lithium nickel oxide (LiNiO2) 12031-95-7, Lithium titanium oxide (Li4Ti5012) 12042-37-4, Alli 12057-17-9, Lithium manganese oxide (LiMn204) 12057-22-6, Lizn 12057-30-6 12162-79-7, Lithium manganese oxide limno2 12190-79-3, Cobalt lithium oxide (CoLiO2) 12332-29-5, Iron lithium nitride (FeLi3N2) 12338-02-2 13843-81-7, DiLithium dichromate 14024-11-4, Lithium tetrachloroaluminate 14283-07-9. Lithium tetrafluoroborate 14307-35-8, Lithium chromate 15365-14-7, Iron lithium phosphate felipo4 18424-17-4, Lithium hexafluoroantimonate 21324-40-3, Lithium hexafluorophosphate 25743-90-2 Lithium hexafluoroarsenate 33454-82-9, Lithium triflate 35363-40-7, Ethyl propyl carbonate 56525-42-9, Methyl propyl carbonate 61234-06-8, Lithium 80, silicon 20 atomic 62852-65-7, DiLithium decachlorodecaborate(2-) 82906-17-0 97037-11-1 97037-12-2 128975-24-6, LIthium manganese nickel oxide LiMn0.5Ni0.502 135573-53-4, Cobalt lithium nickel oxide co0-llini0-lo2 174421-80-8, Cobalt lithium nitride (Co0.4Li2.6N) 177997-11-4, Cobalt gallium lithium nickel oxide 177997-12-5, Boron cobalt lithium nickel oxide 177997-13-6, Aluminum cobalt lithium nickel oxide 180997-14-2, Cobalt lithium magnesium nickel oxide 184912-51-4, Copper lithium nitride (Cu0.4Li2.6N) 244304-18-5, Cobalt lithium nickel silicon oxide 244304-20-9, Cobalt lithium nickel titanium oxide 244761-29-3, Lithium bisoxalatoborate 291298-96-9 321201-33-6, Lithium tris(oxalato)phosphate 346417-97-8, Cobalt lithium manganese nickel oxide (Co0.33LiMn0.33Ni0.33O2) 383187-24-4 427879-42-3, Lithium bis(difluoromalonato)borate 471294-34-5 476300-71-7, Lithium carbide (LiC6) 913080-19-0, Lithium (difluoromalonato) (oxalato)borate 913080-20-3, Lithium tris(difluoromalonato) phosphate RL: TEM (Technical or engineered material use); USES (Uses)

(triazine compds. for removing acids and water from nonaq. electrolytes for electrochem. cells)

L35 ANSWER 6 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN AN 2006:1256604 HCAPLUS Full-text

DN 146:30076

TI Nonaqueous electrolytic solution for electrochemicals cells

IN Xu, Wu; Deng, Zhongvi; Bolomev, Pascal

PA Ferro Corporation, USA

SO U.S. Pat. Appl. Publ., 10pp.

CODEN: USXXCO

DT Patent

LA English

FAN CNT 1

-	I THIN . CIVI I				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
Ι	PI US 20060269846	A1	20061130	US 2005-138905	
					200505
					26

PRAI US 2005-138905

20050526

OS MARPAT 146:30076

AB The invention relates to the use of a nitrogen silylated compound as additive in a nonaq. electrolytic solution The electrolytic solution is suitable for use in electrochem. cells such as lithium and lithium ion batteries. Batteries using this electrolytic solution have long cycle life and high capacity retention.

IT 126-73-8, Tributyl phosphate, uses 15365-14-7,

Iron lithium phosphate felipo4

RL: TEM (Technical or engineered material use); USES (Uses)

(nonaq. electrolytic solution for electrochems. cells)

RN 126-73-8 HCAPLUS

CN Phosphoric acid tributyl ester (CA INDEX NAME)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

Fe(II)

● Tri

CC

ΙT

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INCL 429326000: 429339000: 429336000: 429337000: 429328000: 429329000:
    029623200: 029623300: 029623500
    52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
    57-57-8, β-Propiolactone 68-12-2, Dmf, uses 75-05-8,
    Acetonitrile, uses 78-40-0, Triethyl phosphate
                                                     79-20-9, Methvl
    acetate 96-47-9, 2-Methyltetrahydrofuran 96-48-0,
    γ-Butyrolactone 96-49-1, Ethylene carbonate 105-37-3.
    Ethyl propionate 105-54-4, Ethyl butyrate 105-58-8, Diethyl
               105-66-8, Propyl butyrate 106-36-5, Propyl propionate
    carbonate
    107-31-3, Methyl formate 108-29-2, 4-Methyly-Butyrolactone
    108-32-7, Propylene carbonate 109-21-7, Butyl butyrate 109-60-4,
    Propyl acetate 109-94-4, Ethyl formate 109-99-9, Thf, uses
    110-71-4, 1,2-Dimethoxyethane 110-74-7, Propyl formate 112-48-1,
    1,2-Dibutoxyethane 115-86-6, Triphenyl phosphate 115-96-8,
    Tris(2-chloroethyl)phosphate 123-86-4, Butyl acetate 123-91-1,
    1,4-Dioxane, uses 126-73-8, Tributyl phosphate, uses
    141-78-6, Ethyl acetate, uses 358-63-4,
    Tris(2,2,2-trifluoroethyl)phosphate 512-56-1, Trimethyl phosphate
    513-02-0, Triisopropyl phosphate 513-08-6, Tripropyl phosphate
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    3-Methyly-Butyrolactone 2528-39-4, Trihexyl phosphate
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    7440-44-0, Carbon, uses 7791-03-9, Lithium perchlorate
    12022-46-7. Iron lithium oxide (FeLiO2) 12031-65-1. Lithium nickel
    oxide (LiNiO2) 12031-95-7, Lithium titanium oxide (Li4Ti5012)
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12042-37-4, Alli 12057-17-9, Lithium manganese oxide (LiMn204) 12057-22-6, Lizn 12057-30-6 12162-79-7, Lithium manganese oxide limno2 12190-79-3, Cobalt lithium oxide (CoLiO2) 12332-29-5. Iron lithium nitride (FeLi3N2) 12338-02-2 13843-81-7. Dilithium dichromate 14024-11-4, Lithium tetrachloroaluminate 14283-07-9, Lithium tetrafluoroborate 14307-35-8, Lithium chromate 15365-14-7, Iron lithium phosphate felipo4 18424-17-4, Lithium hexafluoroantimonate 21324-40-3, Lithium hexafluorophosphate 25743-90-2 29935-35-1, Lithium hexafluoroarsenate 33454-82-9, Lithium triflate 35363-40-7, Ethyl propyl carbonate 56525-42-9, Methyl propyl carbonate 61234-06-8, Lithium 80, silicon 20 atomic 62852-65-7, Dilithium decachlorodecaborate(2-) 82906-17-0 97037-11-1 97037-12-2 128975-24-6, Lithium manganese nickel oxide limn0.5ni0.5o2 135573-53-4, Cobalt lithium nickel oxide Co0-1LiNi0-102 174421-80-8, Cobalt lithium nitride (Co0.4Li2.6N) 177997-11-4, Cobalt gallium lithium nickel oxide 177997-12-5, Boron cobalt lithium nickel oxide 177997-13-6, Aluminum cobalt lithium nickel oxide 180997-14-2, Cobalt lithium magnesium nickel oxide 184912-51-4, Copper lithium nitride (Cu0.4Li2.6N) 244304-18-5, Cobalt lithium nickel silicon oxide 244304-20-9. Cobalt lithium nickel titanium oxide 244761-29-3, Lithium bisoxalatoborate 291298-96-9 321201-33-6, Lithium tris(oxalato)phosphate(1-) 346417-97-8, Cobalt lithium manganese nickel oxide (Co0.33LiMn0.33Ni0.33O2) 383187-24-4 427879-42-3 471294-34-5 476300-71-7, Lithium carbide (LiC6) 913080-19-0 913080-20-3 RL: TEM (Technical or engineered material use); USES (Uses) (nonag. electrolytic solution for electrochems. cells)

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L35 ANSWER 7 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN
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AN 2006:1124296 HCAPLUS Full-text

DN 145:457702

TΙ Nonagueous electrolytic solution for lithium secondary battery

IN Xu, Wu; Deng, Zhongyi; Bolomey, Pascal

PA Ferro Corporation, USA

SO U.S. Pat. Appl. Publ., 6pp. CODEN: USXXCO

DТ Patent

LA FAN.	English				
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡΙ	US 20060240327	A1	20061026	US 2005-113823	200504 25

US 7255965 B2 20070814

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WO 2006115737 A1 20061102 WO 2006-US13113
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    MARPAT 145:457702
OS
    The use of lithium bis(oxalate)borate as an additive in a lithium
AB
     secondary battery provides improved battery performance such as long
     life and high capacity retention after high temperature storage.
    126-73-8, Tributyl phosphate, uses 15365-14-7,
ΙT
     Iron lithium phosphate felipo4
     RL: DEV (Device component use): USES (Uses)
        (nonag, electrolytic solution for lithium secondary battery)
RN
     126-73-8 HCAPLUS
    Phosphoric acid tributyl ester (CA INDEX NAME)
CN
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RN 15365-14-7 HCAPLUS CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

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INCL 429324000; 429338000; 429342000; 429343000; 429339000; 429326000;
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    57-57-8, β-Propiolactone 68-12-2, Dmf, uses 75-05-8,
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    Acetonitrile, uses 78-40-0, Triethyl phosphate 79-20-9, Methyl
    acetate 96-47-9, 2-Methyl tetrahydrofuran 96-48-0,
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    acetate, uses 358-63-4, Tris(2,2.2-trifluoroethyl)phosphate
    512-56-1, Trimethyl phosphate 513-02-0, Triisopropyl phosphate
    513-08-6, Tripropyl phosphate 542-28-9, 8-Valerolactone
    542-52-9, Dibutyl carbonate 554-12-1, Methyl propionate 590-01-2, Butyl propionate 592-84-7, Butyl formate 616-38-6,
    Dimethyl carbonate 623-42-7, Methyl butyrate 623-53-0, Ethyl
    methyl carbonate 623-96-1, Dipropyl carbonate 629-14-1,
    1,2-Diethoxyethane 646-06-0, 1,3-Dioxolane 1330-78-5, Tritolyl
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L35

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DN TΙ

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DT

US 7238453

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    7440-44-0, Carbon, uses 7791-03-9, Lithium perchlorate
    12022-46-7, Iron lithium oxide (FeLiO2) 12031-65-1, Lithium nickel
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    12042-37-4, Alli 12057-17-9, Lithium manganese oxide (LiMn204)
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    tetrafluoroborate 14307-35-8, Lithium chromate 15365-14-7
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    RL: DEV (Device component use); USES (Uses)
       (nonag. electrolytic solution for lithium secondary battery)
            THERE ARE 38 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE.CNT 38
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
   ANSWER 8 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN
    2006:1124255 HCAPLUS Full-text
    Nonagueous electrolytic solution with mixed salts
IN Xu, Wu; Deng, Zhongvi; Bolomev, Pascal; Pavne, Martin W.
   Ferro Corporation, USA
SO U.S. Pat. Appl. Publ., 6pp.
    CODEN: USXXCO
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LA English
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                      KIND DATE
                                        APPLICATION NO.
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PI US 20060240322 A1
                             20061026 US 2005-113966
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B2 20070703

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US 20060236528
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WO 2006115681
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PRAI	US 2005-111823	A2	20050422		
	US 2005-113823	A	20050425		
	US 2005-113966	A2	20050425		
	US 2005-196782	A	20050803		
	WO 2006-US11539	W	20060329		
	WO 2006-US13113	W	20060410		
OS	MARPAT 145:457700				

AB The use of at least two electrolyte salts in a lithium secondary battery provides improved battery performance such as long cycle life of high discharge capacity and high capacity retention.

IT 126-73-8, Tributyl phosphate, uses 15365-14-7,

Iron lithium phosphate felipo4

RL: DEV (Device component use); USES (Uses)

(nonaq. electrolytic solution with mixed salts)

RN 126-73-8 HCAPLUS

CN Phosphoric acid tributyl ester (CA INDEX NAME)

RN 15365-14-7 HCAPLUS

CN Phosphoric acid, iron(2+) lithium salt (1:1:1) (9CI) (CA INDEX NAME)

● Fe(II)

● Li

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INCL 429188000: 429338000: 429342000: 429343000: 429337000: 429329000:
    429330000; 429331000; 429332000; 429231100
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    Acetonitrile, uses 78-40-0, Triethyl phosphate
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    acetate 96-47-9, 2-Methyltetrahydrofuran 96-48-0,
    γ-Butyrolactone 96-49-1, Ethylene carbonate 105-37-3,
    Ethyl propionate 105-54-4, Ethyl butyrate 105-58-8, Diethyl
    carbonate 105-66-8, Propyl butyrate 106-36-5, Propyl propionate,
    uses 107-31-3, Methyl formate 108-29-2,
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    126-73-8, Tributyl phosphate, uses 141-78-6, Ethyl
    acetate, uses 358-63-4, Tris(2,2,2-trifluoroethyl)phosphate
    512-56-1, Trimethyl phosphate 513-02-0, Triisopropyl phosphate
    513-08-6, Tripropyl phosphate 542-28-9, δ-Valerolactone
    542-52-9, Dibutyl carbonate 554-12-1, Methyl propionate
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    Dimethyl carbonate 623-42-7, MEthyl butyrate 623-53-0, Ethyl
    methyl carbonate 623-96-1, Dipropyl carbonate 629-14-1,
    1,2-Diethoxyethane 646-06-0, 1,3-Dioxolane 1330-78-5, Tritolyl
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    3-Methyl-y-Butyrolactone 2528-39-4, Trihexyl phosphate
    4437-85-8, Butylene carbonate 7439-93-2, Lithium, uses
    7440-44-0, Carbon, uses 7791-03-9, LIthium perchlorate
    12022-46-7, Iron lithium oxide (FeLiO2) 12031-65-1, Lithium nickel
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12042-37-4, Alli 12057-17-9, Lithium manganese oxide (LiMn204)
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    Copper lithium nitride (Cu0.4Li2.6N) 244761-29-3, Lithium
    bisoxalatoborate 291298-96-9 321201-33-6 346417-97-8, Cobalt
    lithium manganese nickel oxide (Co0.33LiMn0.33Ni0.33O2)
    383187-24-4 427879-42-3 476300-71-7, Lithium carbide (LiC6)
    913080-19-0 913080-20-3
    RL: DEV (Device component use); USES (Uses)
       (nonag, electrolytic solution with mixed salts)
RE.CNT 35
            THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
L35 ANSWER 9 OF 9 HCAPLUS COPYRIGHT 2009 ACS on STN
    1999:708588 HCAPLUS Full-text
    131:327534
    Fibrinogen-coated droplets of liquid hydrophobic phases
   Retzinger, Gregory S.; Deanglis, Ashley P.
   University of Cincinnati, USA
SO PCT Int. Appl., 62 pp.
    CODEN: PIXXD2
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FAN.CNT 1
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A1 19991104 CA 1999-2327469

199904

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EP 1073425 A1 20010207 EP 1999-921733

> 199904 28

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT. IE. FI

PRAI US 1998-83709P

Р 19980430

WO 1999-US9940 W 19990428 AB

Fibrinogen adsorbs spontaneously from aqueous media to droplets of liquid hydrophobic (oil) phases dispersed in those same media. Examples of such phases include mineral oils, straight chain hydrocarbons, and various plant- and animal-derived oils. Lecithin pre-existing on the surface of oil droplets reduces significantly the amount of fibrinogen that can otherwise bind to them. When bound, fibringen remains active in the classic sense of fibrin gelation. As a consequence, oil droplets coated with fibringen can participate in biol. important adhesive processes in which the protein would be expected to participate. Certain polyanions, e.g., heparin, pentosan polysulfate, dextran sulfate and suramin, bind to adsorbed fibrin(ogen) and prevent thrombin-dependent adhesion of fibrinogencoated surfaces. Thus, these polyanions can be used to prevent adhesion between fibrin(ogen)-coated oil droplets and other fibrin(ogen)-coated surfaces. Potential practical applications and biol. implications of these phenomena include treating or assessing diseases associated with fibrin clots, other fibrin(ogen)-coated surfaces, and/or fibrin(ogen)-associated processes, and for formulation of vehicle for the targeted delivery of drugs and as an adjuvant for vaccines.

112-40-3, Dodecane TT

RL: PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)

(fibrin(ogen)-coated droplets of hydrophobic phases for diagnostic and therapeutic uses)

112-40-3 HCAPLUS RN

CN Dodecane (CA INDEX NAME)

Me- (CH2) 10-Me

IΤ 140207-93-8, Sodium pentosan polysulfate RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(polyanions prevention of adhesion between fibrin(ogen)-coated

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oil droplets and other fibrin(ogen)-coated surfaces)
RN
    140207-93-8 HCAPLUS
CN
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     sodium salt (CA INDEX NAME)
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     CRN 9062-57-1
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    CCI PMS, MAN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
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    CMF H2 O4 S
    ICM A61K009-50
T.C.
    ICS A61K009-16
    63-6 (Pharmaceuticals)
CC
    111-01-3, Squalane 111-02-4, Squalene 112-40-3, Dodecane
ΙT
    RL: PEP (Physical, engineering or chemical process); THU
     (Therapeutic use); BIOL (Biological study); PROC (Process); USES
     (Uses)
        (fibrin(ogen)-coated droplets of hydrophobic phases for
        diagnostic and therapeutic uses)
    129-46-4, Sodium suramin 9041-08-1, Heparin sodium 9042-14-2,
ΙT
     Dextran sulfate 140207-93-8, Sodium pentosan polysulfate
     157009-81-9
     RL: PRP (Properties); THU (Therapeutic use); BIOL (Biological
     study); USES (Uses)
        (polyanions prevention of adhesion between fibrin(ogen)-coated
        oil droplets and other fibrin(ogen)-coated surfaces)
RE.CNT 1
              THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS RECORD
              ALL CITATIONS AVAILABLE IN THE RE FORMAT
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